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(54) Title: IMMUNOLIPOSOMES THAT OPTIMIZE INTERNALIZATION INTO TARGET CELLS				
(57) Abstract <p>The present invention provides for immunoliposomes that optimizes internalization of a drug into target cells bearing a characteristic cell surface marker. The immunoliposomes comprise an Fab' domain of an antibody that specifically binds the characteristic marker, an amphipathic vesicle-forming lipid, and a polyethylene glycol derivatized lipid. The invention also provides for growth-inhibiting immunoliposomes that lack growth-inhibiting therapeutic agents and yet are capable of inhibiting the growth and proliferation of target cells.</p>				

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# IMMUNOLIPOSOMES THAT OPTIMIZE INTERNALIZATION INTO TARGET CELLS

## FIELD OF THE INVENTION

The present invention relates to the field of liposomes. In particular,  
5 the present invention relates to liposomes specifically targeted to characteristic  
markers on target cells and which contain up to 4 mole percent of a hydrophilic  
polymer which results in an unexpected high rate of cellular incorporation.

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10 Institutes of Health. The Government of the United States has certain rights in this  
invention.

## BACKGROUND OF THE INVENTION

A number of pharmaceutical agents and potential pharmaceutical  
15 agents suffer from poor aqueous solubility, high levels of antigenicity, toxicity, or  
rapid degradation in serum which can hamper the development of suitable clinical  
formulations. One solution to these problems has been to encapsulate the  
pharmaceutical agent in a delivery vehicle that is soluble in aqueous solutions and  
that shields the agent from direct contact with tissues and blood. In particular,  
20 formulations based on liposome technology are of significant interest. Liposomes  
are vesicles comprised of concentrically ordered phospholipid bilayers which  
encapsulate an aqueous phase. They form spontaneously when phospholipids are  
exposed to aqueous solutions and can accommodate a variety of bioactive  
molecules.

25 Liposomes have proved a valuable tool as an *in vivo* delivery system  
for enhancing the efficacy of various pharmacologically active molecules (Ostro et  
al. *Liposomes from Biophysics to Therapeutics*, Dekker, New York, pp. 1-369  
(1987)). Animal studies have shown that liposomes can decrease the toxicity of  
several antitumor and antifungal drugs, leading to clinical trials with promising  
30 results (Sculier et al. *Eur. J. Cancer Clin. Oncol.*, 24: 527-538; Gabizon, et al.

Eur. J. Cancer Clin. Oncol., 25: 1795-1803 (1989); Treat *et al.*, *J. Natl. Cancer Inst.*, 82: 1706-1710 (1990); Lopez-Berestein *et al.* *J. Infect. Dis.*, 151: 704-710 (1985); Presant *et al.* *Cancer*, 62: 905-911 (1988)). In addition, liposomes have been shown to be efficient carriers of antiparasitic drugs for treating intracellular  
5 infections of the reticuloendothelial system (RES), in activating macrophage cells to become tumoricidal, in models of metastasis, and in enhancing the immune response to encapsulated antigens, thus facilitating the formulation of artificial vaccines (*Liposomes in the Therapy of Infectious Diseases and Cancer*, Lopez-Berestein & Fidler, eds. Liss, New York (1989); Alving *et al.* *Immunol. Lett.*, 25:  
10 275-280 (1990)).

All these effects stem from the capacity of macrophage cells in the liver and spleen (mononuclear phagocytic system MPS or reticuloendothelial system RES) to remove the majority of liposomes from the blood circulation within minutes (*Liposomes as Drug Carriers*, Gregoriadis, ed., Wiley, New York.  
15 (1988)). Such rapid clearance of liposomes however, has limited their prospects as an *in vivo* delivery system for transporting drugs to sites of disease beyond the RES.

Recent reports have described the use of various polymers to increase serum half-life of liposomes. In particular, it has been recognized that  
20 formulations of liposomes containing either mono-sialoganglioside (GM<sub>1</sub>) or lipid derivatives of polyethylene glycol avoid MPS removal and significantly increase serum half-life (Allen *et al.* *FEBS Lett.*, 223: 42-46 (1987); Klivanov *et al.*, *FEBS Lett.*, 268: 235-237 (1990); Blume *et al.* *Biochim. Biophys. Acta.*, 1029: 91-97 (1990); Allen *et al.* *Biochim. Biophys. Acta*, 1066: 29-36 (1991); Papahadjopoulos  
25 *et al.*, *Proc. Natl. Acad. Sci. USA*, 88: 11460-11464 (1991); Senior *et al.* *Biochim. Biophys. Acta.*, 1062: 77-82 (1991); Allen *et al.* *Biochim. Biophys. Acta.*, 1068: 133-141 (1991)).

Many reports have demonstrated that rapid removal of circulating liposomes *in vivo* by cells of the mononuclear phagocytic system (MPS) can be  
30 overcome by incorporation of lipids derivatized with the hydrophilic polymer polyethylene glycol (PEG). These liposomes are referred to as sterically stabilized

or "stealth" liposomes. With PEG having a molecular weight in the range of 1000 to 5000, prolonged circulation and reduced MPS uptake is achieved (Woodle & Lasic. *Biochim. Biophys. Acta* 1113:171-199 (1992)). However, this reduction in clearance by the MPS is also associated with a reduction in uptake by a variety of cells (Lee, K.D. et al. *Biochim. Biophys. Acta* 1103:185-197 (1992)). In addition, the presence of hydrophilic polymers on the surface of the liposome appears to interfere with specific ligand recognition by targeting moieties conjugated to the liposome. Presumably this occurs due to steric hinderance of the active site of the targeting moiety by the long chain PEG molecules. (Klibanov et al. *Biochim. Biophys. Acta* 1062:148-148 (1991)).

Finally, while most therapeutic agents transported by liposomes must enter the cytoplasm of the target cell in order to express their biological activity, it is generally appreciated that most liposomes are either not actually internalized by the target cells, or, where uptake does occur, it is generally via an endocytotic pathway. Thus actual drug to the target cell typically entails release from the liposome (e.g. through disruption of the liposome itself or through "leakage") in the vicinity of the target cell and then subsequent uptake (either through diffusion, endocytosis, phagocytosis, or active transport) of the therapeutic agent from solution by the target cell. Indeed immunoliposomes have been designed to actually induce destabilization and fragmentation of the liposome once the targeting antibody has bound a target, thereby freeing the liposome contents (see, U.S. Patent No. 4,957,735). Even these "target-sensitive" liposomes, lose a considerable amount of the therapeutic agent in solution before it can be taken up by the target cell. Alternatively, if the liposome is internalized by an endocytotic process, it is ultimately incorporated in a lysosome where strong acid conditions exist that can degrade a number of therapeutic agents (e.g. proteins).

Thus, delivery of effective doses of therapeutic agents to the cytoplasm of the target cell is hampered by low residence times in serum, ineffective targeting when residence times are increased, considerable loss of the therapeutic agent in solution before it may be taken up by the target cell, and degradation of the therapeutic in the endosomic/lysosomal pathway. Clearly, it

would be desirable to obtain a liposome with increased serum half-life, capable of specifically targeting particular cells, and also capable of being internalized into the cytoplasm by the target cells thereby avoiding loss of the therapeutic agent or degradation by the endosomal/lysosomal pathway.

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### SUMMARY OF THE INVENTION

The present invention provides novel immunoliposomes optimized for delivering therapeutic agents to the cytoplasm of a target cell. These immunoliposomes exhibit increased half-life in blood, are capable of specifically  
10 targeting particular cells, and are capable of being internalized into the cytoplasm by the target cells thereby avoiding loss of the therapeutic agents or degradation by the endolysosomal pathway.

Thus, in one preferred embodiment this invention provides for immunoliposomes that optimize internalization of a therapeutic agent into the  
15 cytoplasm of a cell bearing a characteristic cell surface marker. These immunoliposomes comprise an Fab' domain of an antibody wherein the Fab' domain specifically binds the characteristic marker, an amphipathic vesicle-forming lipid that forms a liposome, a polyethylene glycol derivatized lipid wherein the polyethylene glycol has an average molecular weight of between about 750 D and  
20 5000 D, more preferably between about 1200 D and about 3000 D, most preferably about 1900 D, and a therapeutic agent contained within the liposome. The derivatized lipid is present at up to about 1.2 mole percent, more preferably at up to about 2.4 mole percent, and most preferably at up to about 3.6 mole percent of total lipid. Preferred characteristic markers include growth factor  
25 receptors. Particularly preferred are growth factor receptors including HER1, HER2, HER3 and HER4 with HER2 being most preferred. The Fab' domain may be a humanized Fab' domain, more specifically a humanized Fab' domain of an anti-HER2 monoclonal antibody. The growth-inhibiting immunoliposome may further comprise a maleimide-derivatized phosphatidylethanolamine (M-PE) which  
30 forms a thioether linkage to the Fab' domain of an antibody. The vesicle forming lipid may include a phospholipid, a glycolipid, a sphingolipid, or a sterol. The

immunoliposomes have an average diameter that ranges from about 50 nm to about 500 nm, more preferably about 75 nm to about 300 nm and most preferably is about 100 nm. Therapeutic agents in the liposome may include daunomycin, idarubicin, mitoxantrone, mitomycin, cisplatin and other Platinum II analogs, vincristine, epirubicin, aclacinomycin, methotrexate, etoposide, doxorubicin, cytosine arabinoside, fluorouracil and other fluorinated pyrimidines, purines, or nucleosides, bleomycin, mitomycin, plicamycin, dactinomycin, cyclophosphamide and derivatives thereof, thiotepa, BCNU, taxol, taxotere and other taxane derivatives and isolates, camptothecins, polypeptides, a nucleic acid, a nucleic acid having a phosphorothioate internucleotide linkage, and a nucleic acid having a polyamide internucleotide linkage.

In one particularly preferred immunoliposome, the antibody Fab' domain is that of rhuMAbHER2, with the Fab' domain conjugated to maleimide-derivatized phosphatidylethanolamine (M-PE), the vesicle forming lipid is phosphatidylcholine (PC) and cholesterol (Chol) and the polyethylene glycol derivatized lipid is polyethylene glycol derivatized phosphatidylethanolamine (PEG-PE) where the polyethylene glycol component has a molecular weight of about 1900 D, and where the ratio PC:Chol:M-PE is 150:100:3 and the PEG-PE is present in an amount up to about 3.6 mole percent of total lipid.

This invention also provides for a method of optimizing internalization of a therapeutic agent into a cell bearing a characteristic marker, the method comprising contacting the cell with any of the immunoliposomes summarized above and internalizing the liposome contents into the cytoplasm of the cell. This internalization may be by fusion of the liposome with cell membranes, or early exit from the endocytic vesicles.

In another preferred embodiment, this invention provides for a growth-inhibiting immunoliposome that specifically binds to a cell bearing a characteristic cell surface marker thereby inhibiting the proliferation or growth of that cell. The immunoliposome comprises an Fab' domain of an antibody where the Fab' domain specifically binds the marker and an amphipathic vesicle-forming lipid that forms a liposome. Unlike typical drug-delivery liposomes, however, this

liposome contains no growth-inhibiting therapeutic agent and may contain no therapeutic agent at all. A composition may be identified as not being a growth-inhibiting therapeutic agent when the decrease in proliferation rate of target cells is less than 10 percent, more preferably less than 5 percent, more than the decrease in proliferation rate of the target cells caused by an "empty" liposome containing no agent (other than a neutral buffer or water) at all. Preferred characteristic markers include growth factor receptors. Particularly preferred are growth factor receptors including HER1, HER2, HER3 and HER4 with HER2 being most preferred. The Fab' domain may be a humanized Fab' domain, more specifically a humanized Fab' domain of an anti-HER2 (anti-p185HER2) monoclonal antibody. The growth-inhibiting immunoliposome may further comprise a maleimide derivatized phosphatidylethanolamine (M-PE) which forms a thioether linkage to the Fab' domain of an antibody. The vesicle forming lipid may include a phospholipid, a glycolipid, a sphingolipid, or a sterol. The growth-inhibiting immunoliposome may also comprise a hydrophilic polymer. Preferred hydrophilic polymers include polyethylene glycol, polypropylene glycol, mono-sialoganglioside (GM<sub>1</sub>), phosphatidylinositol (PI) or cerebroside sulfate (CS)). Polyethylene glycol, when incorporated into the lipid will be incorporated as a polyethylene glycol derivatized lipid, preferably a polyethylene glycol derivatized phospholipid such as PEG-PE. The molecular weight of the polyethylene glycol may range from about 750 D to about 5000 D, more preferably from about 1200 D to about 3000 D, and most preferably is about 1900 D. The liposomes have an average diameter that ranges from about 50 nm to about 500 nm, more preferably about 75 nm to about 300 nm and most preferably is about 100 nm. In one particularly preferred immunoliposome, the antibody Fab' domain is rhuMAbHER2, the Fab' domain is conjugated to M-PE, the vesicle forming lipid is phosphatidylcholine (PC) and cholesterol (Chol) and the polyethylene glycol derivatized lipid is polyethylene glycol derivatized phosphatidylethanolamine (PEG-PE) where the polyethylene glycol component has a molecular weight of about 1900 D, and where the ratio PC:Chol:M-PE is 150:100:3 and the PEG-PE is present in an amount up to about 3.6 mole percent of total lipid.



This invention also provides for a method of inhibiting growth of a cell bearing a characteristic marker, the method comprising contacting the cell with any of the growth-inhibiting liposomes summarized above.

In yet another embodiment, this invention also provides for pharmaceutical compositions comprising any of the growth-inhibiting or therapeutic-agent carrying immunoliposomes described above. The pharmaceutical compositions comprise a therapeutically effective dose of the immunoliposome and a pharmaceutically acceptable carrier or excipient.

#### 10 **BRIEF DESCRIPTION OF THE DRAWINGS**

Figure 1 illustrates flow cytometric histograms showing binding of anti-p185<sup>HER2</sup> immunoliposomes to SK-BR-3 cells. Immunoliposomes bound to SK-BR-3 cells after washing were detected by FITC-labeled goat anti-human IgG, which recognizes rhuMabHER2 Fab' fragments. SK-BR-3 cells were incubated with conventional immunoliposomes (A), sterically stabilized (6 mole% PEG-PE) immunoliposomes (B), and free rhuMabHER2-Fab' fragments (C) at equivalent antibody concentrations (3.3 µg/ml).

Figure 2 shows the binding of anti-p185<sup>HER2</sup> conventional immunoliposomes to BT-474 cells. (A) BT-474 cells in monolayer culture were treated with conventional immunoliposomes in the presence of competing <sup>125</sup>I-labeled rhuMabHER2 as described in Methods. (B) Scatchard transformation of the data shown in (A).

Figure 3 illustrates the antiproliferative activity of anti-p185<sup>HER2</sup> immunoliposomes against SK-BR-3 cells. SK-BR-3 cells in monolayer culture were treated with immunoliposomes at antibody doses indicated on the abscissa, and relative cell proliferation determined as described in Methods. Control liposomes lacking antibody were dosed according to liposome concentration, and are plotted at equivalent liposome concentration with the appropriately matched immunoliposome. Control (no antibody) conventional liposomes; control sterically stabilized (6 mole% PEG-PE) liposomes; conventional immunoliposomes; anti-p185<sup>HER2</sup> sterically stabilized (6 mole% PEG-PE) immunoliposomes; free

(non-liposomal) rhuMAbHER2-Fab'; and free rhuMAbHER2 bivalent antibody are shown as indicated in the key.

Figure 4 shows the cytotoxicity of anti-p185<sup>HER2</sup> immunoliposomes containing doxorubicin. HSPC: Chol immunoliposomes were loaded with doxorubicin as described in Methods. (A) SK-BR-3 cells. (B) WI-38 cells. Cells in culture were treated for 1 hour with: conventional immunoliposomes (triangles); sterically stabilized (2 mole% PEG-PE) immunoliposomes (closed circles); control (irrelevant antibody) sterically stabilized (2 mole% PEG-PE) immunoliposomes (open circles); or free doxorubicin alone (closed squares). Immunoliposomes contained 60-70  $\mu\text{g}$  antibody/ $\mu\text{mol}$  phospholipid and 55-80  $\mu\text{g}$  doxorubicin/ $\mu\text{mol}$  phospholipid; antibody/doxorubicin ratio was 0.8-1.2. Cells were counted 3 days after treatment as described in Methods.

## **DESCRIPTION OF THE PREFERRED EMBODIMENT**

### **15 Definitions and General Parameters**

The following definitions are set forth to illustrate and define the meaning and scope of the various terms used to describe the invention herein.

The following abbreviations are used herein: DOX, doxorubicin; Chol, cholesterol; PA, phosphatidic acid; PC, phosphatidylcholine; PI, phosphatidylinositol; SM, sphingomyelin; M-DPE, maleimide derivatized dipalmitoylethanolamine; PBS, phosphate buffered saline; LUV, large unilamellar vesicles; MLV, multilamellar vesicles; PE, phosphatidylethanolamine; PEG, polyethylene glycol; PEG-PE, polyethylene glycol derivatized phosphatidylethanolamine.

25 The term "amphipathic vesicle-forming lipid" is intended to include any amphipathic lipid having hydrophobic and polar head group moieties, and which by itself can form spontaneously into bilayer vesicles in water, as exemplified by phospholipids, or (b) is stably incorporated into lipid bilayers in combination with phospholipids with its hydrophobic moiety in contact with the interior, hydrophobic region of the bilayer membrane, and its polar head group moiety oriented toward the exterior, polar surface of the membrane. An example

30

of the latter type of vesicle-forming lipid is cholesterol and cholesterol derivatives such as cholesterol sulfate and cholesterol hemisuccinate.

As used herein, the term "specific binding" refers to that binding which occurs between such paired species as enzyme/substrate, receptor/agonist, antibody/antigen, and lectin/carbohydrate which may be mediated by covalent or non-covalent interactions or a combination of covalent and non-covalent interactions. When the interaction of the two species produces a non-covalently bound complex, the binding which occurs is typically electrostatic, hydrogen-bonding, or the result of lipophilic interactions. Accordingly, "specific binding" occurs between a paired species where there is interaction between the two which produces a bound complex having the characteristics of an antibody/antigen or enzyme/substrate interaction. In particular, the specific binding is characterized by the binding of one member of a pair to a particular species and to no other species within the family of compounds to which the corresponding member of the binding member belongs. Thus, for example, an antibody preferably binds to a single epitope and to no other epitope within the family of proteins. The terms "ligand" or "targeting moiety", as used herein, refer generally to all molecules capable of specifically binding to a particular target molecule and forming a bound complex as described above. Thus the ligand and its corresponding target molecule form a specific binding pair. Examples include, but are not limited to antibodies, lymphokines, cytokines, receptor proteins such as CD4 and CD8, solubilized receptor proteins such as soluble CD4, hormones, growth factors, and the like which specifically bind desired target cells, and nucleic acids which bind corresponding nucleic acids through base pair complementarity. Particularly preferred targeting moieties include antibodies and antibody fragments (*e.g.*, the Fab' domain).

The term "immunoliposome" refers to a liposome bearing an antibody or antibody fragment that acts as a targeting moiety enabling the liposome to specifically bind to a particular "target" molecule that may exist in solution or may be bound to the surface of a cell. Where the target molecule is one that is typically found in relative excess (*e.g.*,  $\geq 10$ -fold) and in association with a

particular cell type or alternatively in a multiplicity of cell types all expressing a particular physiological condition the target molecule is said to be a "characteristic marker" of that cell type or that physiological condition. Thus, for example, a cancer may be characterized by the overexpression of a particular marker such as the HER2 (*c-erbB-2/neu*) proto-oncogene in the case of breast cancer.

A "hydrophilic polymer" as used herein refers to long chain highly hydrated flexible neutral polymers attached to lipid molecules. Examples include, but are not limited to polyethylene glycol-, or polypropylene glycol-modified lipids PI or CS, or ganglioside GM<sub>1</sub>.

The term "mole percent" when referring to the percentage of hydrophilic polymer in a liposome is expressed relative to the total lipid in the liposome unless otherwise stated. Thus, for example, in a liposome comprising a ratio of phosphatidylcholine (PC) to cholesterol (Chol) of 150:100, a 4 mole percent of hydrophilic polymer (*e.g.* PEG) would represent a ratio of PC:Chol:PEG of about 150:100:10.

The term "proliferation" refers to cell division or mitosis. Proliferation may be measured by standard assays such as by uptake of radioactive nucleotides (thymidine) or by direct observation.

## **Immunoliposomes In Drug Delivery**

In one embodiment, the present invention provides immunoliposomes for selective delivery of therapeutic agents to specific tissues in a host and methods of use for those liposomes. The liposomes of this invention employ a composition that optimizes internalization of the liposome into the cytoplasm of the cells of the target tissue. The phrase "optimizes internalization" or "optimal internalization" is used to refer to the delivery of liposome contents such that it achieves maximum delivery to the cytoplasm of the target cell and therefore maximum therapeutic effect. It is recognized that internalization of an immunoliposome into the cytoplasm of a cell is a function of the blood half-life of the liposome, the ability of the liposome to recognize and bind to its target cell, and the uptake of the liposome into the cytoplasm of the target cell. It is well known that addition of a

hydrophilic polymer to liposomes increases serum half-life by decreasing both liposome agglomeration (aggregation) and liposome uptake by the RES. Without being bound to a theory, it is believed that hydrophilic polymers at high concentrations interfere with recognition and binding by the targeting moiety or  
5 ligand and with subsequent uptake by the target cell, thereby decreasing the internalization of the liposome contents by the target cell. Optimal internalization into the cytoplasm of the cell refers to that condition in which maximal uptake into the cytoplasm of the target cell is achieved while still maintaining a blood half-life significantly greater than the blood half-life of liposomes lacking any hydrophilic  
10 polymer and adequate for targeting purposes.

In particular, this invention relies, in part, on the unexpected discovery that a liposome comprising a hydrophilic polymer (*e.g.*, PEG-modified lipid) in an amount up to about 3.6 mole percent of total (vesicle-forming) lipid demonstrates an unexpected high rate of internalization into the cytoplasm of the  
15 target cell while retaining a blood half-life substantially greater than that seen in liposomes lacking a hydrophilic polymer.

In addition, it was also an unexpected discovery that when liposomes comprising up to 3.6 mole percent of a hydrophilic polymer were conjugated with an Fab' fragment of an antibody as a targeting moiety the liposome shows a high  
20 degree of cellular specificity and a binding affinity greater than that of the Fab' fragments alone. In fact, the binding specificity achieved by the immunoliposomes of the present invention is comparable to the binding specificity of the intact antibody. This result is particularly surprising since the intact antibody is a tetramer that comprises a pair of variable domain "arms" that are largely  
25 responsible for antibody specificity and avidity. Fab' regions, consisting of only one "arm" typically lack the specificity and binding avidity of the intact antibody. Thus they are typically expected to form poor targeting moieties.

The immunoliposomes of this invention optimize delivery of therapeutic agents to the cytoplasm of the target cell by maintaining an elevated  
30 blood half-life, as compared to a liposome lacking a hydrophilic polymer, by maintaining a high degree of target specificity, and by effective internalization of

the liposome itself (carrying therapeutic agent) thereby avoiding considerable loss of the therapeutic agent in solution or degradation of the therapeutic in the endosomic/lysosomal pathway. The liposomes of the present invention are thus particularly useful as vehicles for the delivery of therapeutics to specific target  
5 cells.

### Immunoliposomes as Cell Growth Inhibitors

This invention also provides for growth-inhibiting immunoliposomes that may be utilized to inhibit tumor cell proliferation and thus provide an  
10 antitumor activity without encapsulating a growth-inhibiting therapeutic agent. In fact, the growth-inhibiting immunoliposomes of the present invention are effective when they contain no therapeutic agent. The growth-inhibiting immunoliposomes of this invention generally comprise an Fab' domain of an antibody that specifically binds to a cell bearing a characteristic marker, and an amphipathic vesicle forming  
15 lipid. In particular, the liposome may be conjugated to the Fab' domain of an anti-HER2 monoclonal antibody. In a preferred embodiment, the antibody is the Fab' fragment of the human monoclonal anti-HER2 antibody (rhuMAbHER2-Fab'). Unlike free rhuMAbHer2-Fab' in solution, liposomal (membrane) anchoring of the monovalent Fab' fragment results in antiproliferative and antitumor activity  
20 comparable to bivalent rhuMAbHER2. The antibody rhuMAbHER2-Fab' in solution does not have this property. Without being bound to a particular theory, it is believed that membrane anchoring of the Fab' fragment in the anti-HER2 immunoliposome confers this antiproliferative property presumably by enabling cross-linking of p185<sup>HER2</sup> on the tumor cell surface.

25 As indicated above, the growth-inhibiting immunoliposomes do not contain a growth-inhibiting agent. A "growth-inhibiting agent" refers to a chemical agent that reduces the growth rate of cells to which it is administered. In the extreme case a growth-inhibiting agent may be cytotoxic to the cell to which it is administered. As used herein the growth rate of cells refers to the rate of  
30 proliferation of the cells. Increased proliferation rate is typically associated with increased metabolic rate and thus proliferation rates may be assayed by detecting

metabolic rates (*e.g.*, by uptake of a labeled metabolic precursor such as tritiated thymidine). Thus, increased growth or proliferation rate may be taken as indicating an increased metabolic rate or vice versa.

Growth-inhibiting agents are well known to those of skill in the art and include, but are not limited to doxorubicin, ricin A, gelonin. It will be recognized that some compositions (*e.g.*, antibiotics) may exhibit minor growth inhibitory activity as an incidental consequence of their primary activity. Such compositions are not considered herein to be growth-inhibiting agents. The phrase a "liposome containing no growth-inhibiting therapeutic agent" is intended to capture the fact that the inhibition of cell growth and proliferation obtained with the growth-inhibiting immunoliposomes of the present invention is a consequence of the liposome/Fab' construct itself and is not a consequence of the liposome contents. Thus, a growth inhibiting agent, as used herein refers to an agent that, when present in the growth inhibiting immunoliposome, results in a decrease in cell proliferation rate at least 10 percent greater than the decrease in cell proliferation rate observed by administration of the same immunoliposomes lacking any therapeutic or growth inhibiting agent.

While the growth-inhibiting liposomes of this invention will inhibit cell growth and proliferation even when they carry no therapeutic agent and therefore may be administered "empty", one of skill will appreciate that it may be desirable to encapsulate a therapeutic agent other than a growth-inhibiting therapeutic agent thereby achieving a liposome that shows dual, additive or supradadditive activities. Thus, for example, an immunoliposome loaded with an antibiotic will show both antibiotic activity as well as the ability to inhibit growth and proliferation of the target cells.

The growth-inhibiting liposomes and the therapeutic agent-carrying immunoliposomes of the present invention may be utilized to inhibit tumor cell proliferation or to target therapeutics to specific cells in a wide variety of hosts. Preferred hosts include mammalian species such as humans, non-human primates, dogs, cats, cattle horses, sheep, rodents, largomorphs and the like.

### Liposome Composition

In general, the immunoliposomes of the present invention comprise one or more vesicle-forming lipids, an Fab' domain of an antibody which acts as a targeting moiety and, especially in the case of the therapeutic agent delivering  
5 immunoliposomes, a hydrophilic polymer. Without being bound to a particular theory the vesicle-forming lipids act to form a bilayer which encapsulates the therapeutic agent when it is present, the hydrophilic polymer serves to prevent agglomeration of the liposomes and also to decrease uptake of the liposome by the RES and thereby increase blood half-life, and the ligand serves to specifically bind  
10 the liposomes to a cell or tissue bearing a target (*i.e.*, a characteristic marker) for which the ligand is specific. The low mole percentage of the hydrophilic polymer coupled with the use of the Fab' antibody fragment allows specific targeting of the liposome and unexpectedly results in a high level of internalization of the entire liposome into the cytoplasm of the target cell.

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#### A) Vesicle-Forming lipids

The vesicle-forming lipid is preferably one having two hydrocarbon chains, typically acyl chains and a polar head group. Included in this class are the phospholipids, such as phosphatidylcholine (PC), phosphatidylethanolamine (PE),  
20 phosphatidic acid (PA), phosphatidylinositol (PI) and sphingomyelin (SM), where the two hydrocarbon chains are typically between about 14-22 carbon atoms in length and have varying degrees of unsaturation or 14-18 carbon chain saturated phospholipids. Also included in this class are the glycolipids such as cerebrosides and gangliosides.

25

In a preferred embodiment, the major lipid component in the liposomes is phosphatidylcholine. Phosphatidylcholines having a variety of acyl chain groups of varying chain length and degree of saturation are available or may be isolated or synthesized by well-known techniques. In general, less saturated phosphatidylcholines are more easily sized, particularly when the liposomes must  
30 be sized below about 0.3 microns, for purposes of filter sterilization. Phosphatidylcholines containing saturated fatty acids with carbon chain lengths in



the range of  $C_{14}$  to  $C_{22}$  are preferred. Phosphatidylcholines with mono or diunsaturated fatty acids and mixtures of saturated and unsaturated fatty acids may also be used. Liposomes useful in the present invention may also be composed of sphingomyelin or phospholipids with head groups other than choline, such as ethanolamine, serine, glycerol and inositol. In particular, phospholipids suitable for formation of liposomes useful in the methods and compositions of the present invention include, *e.g.*, phosphatidylcholine, phosphatidylglycerol, lecithin,  $\beta,\gamma$ -dipalmitoyl- $\alpha$ -lecithin, sphingomyelin, phosphatidylserine, phosphatidic acid, N-(2,3-di(9-(Z)-octadecenyl-oxy))-prop-1-yl-N,N,N-trimethylammonium chloride, phosphatidylethanolamine, lysolecithin, lysophosphatidylethanolamine, phosphatidylinositol, cephalin, cardiolipin, cerebroside, dicetylphosphate, dioleoylphosphatidylcholine, dipalmitoylphosphatidylcholine, dipalmitoylphosphatidylglycerol, dioleoylphosphatidylglycerol, palmitoyl-oleoyl-phosphatidylcholine, di-stearoyl-phosphatidylcholine, stearoyl-palmitoyl-phosphatidylcholine, di-palmitoyl-phosphatidylethanolamine, di-stearoyl-phosphatidylethanolamine, di-myristoyl-phosphatidylserine, di-oleyl-phosphatidylcholine, and the like. Non-phosphorus containing lipids may also be used in the liposomes of the compositions of the present invention. These include, *e.g.*, stearylamine, docetylamine, acetyl palmitate, fatty acid amides, and the like.

Additional lipids suitable for use in the liposomes of the present invention are well known to persons of skill in the art and are cited in a variety of well known sources, *e.g.*, *McCutcheon's Detergents and Emulsifiers* and *McCutcheon's Functional Materials*, Allured Publishing Co., Ridgewood, N.J., both of which are incorporated herein by reference. Preferred liposomes will include a sterol, preferably cholesterol, at molar ratios of from 0.1 to 1.0 (cholesterol:phospholipid). Most preferred liposome compositions are phosphatidylcholine/cholesterol, distearoylphosphatidylcholine/cholesterol, dipalmitoylphosphatidylcholine/cholesterol, and sphingomyelin/cholesterol. Small amounts (*ie.* < 10%) of other derivatized lipids are often present in liposomes having these compositions.

According to one important feature of the invention, the vesicle-forming lipid may be a relatively fluid lipid, typically meaning that the lipid phase has a relatively low liquid to liquid-crystalline melting temperature, *e.g.*, at or below room temperature, or a relatively rigid lipid, meaning that the lipid has a relatively high melting temperature, *e.g.*, up to 60 °C. As a rule, the more rigid, *i.e.*, saturated lipids, contribute to membrane rigidity in a lipid bilayer structure and also contribute to greater bilayer stability in blood. Other lipid components, such as cholesterol, are also known to contribute to membrane rigidity and stability in lipid bilayer structures. As mentioned above, a long chain (*e.g.* C<sub>14</sub>-C<sub>22</sub>) saturated lipid plus cholesterol is one preferred composition for delivering therapeutic compositions to target tissues, such as solid tumors, since these liposomes do not tend to release the drugs into the plasma as they circulate through the blood stream. Phospholipids whose acyl chains have a variety of degrees of saturation can be obtained commercially. For example, egg phosphatidylcholine (EPC) can be purchased from Avanti Polar Lipids (Alabaster, AL) and hydrogenated soy phosphatidylcholine (HSPC) can be obtained from Natterman (Cologne, FRG). Alternatively, phospholipids can be prepared according to published methods. (see D.M. Small, "The physical chemistry of lipids" (1986) Plenum Press, N.Y., or D.D. Lasic, "Liposomes: from physics to applications" (1993) Elsevier, Amsterdam; N.Y.)

#### **B) Hydrophilic Polymer.**

As noted above the presence of hydrophilic polymers tends to increase the blood half-life of a liposome. (See, for example, Woodle, *et al.* *Biochim. Biophys. Acta.*, 1113: 171-199 (1992).) Thus, it is often desirable to add a hydrophilic polymer such as polyethylene glycol (PEG)-modified lipids or ganglioside G<sub>M1</sub> to the liposomes. Addition of such components prevents liposome aggregation during coupling of the targeting moiety to the liposome. These components also provides a means for increasing circulation lifetime of the phospholipid. However, it has been observed that while hydrophilic polymers decrease liposome uptake by the RES and thereby increase blood half-life, there

is a corresponding decrease in uptake by target tissues as well. It was an unexpected discovery of the present invention that a concentration of hydrophilic polymer (*e.g.*, PEG) of 1 to 4 mole percent of vesicle-forming lipid (excluding cholesterol) provides optimal cellular uptake combined with adequate blood half-

5 life.

A number of different methods may be used for the preparation of PEG for incorporation into liposomes. In a preferred embodiment, PEG is incorporated as PEG derivatized phosphatidylethanolamine (PEG-PE) or PEG derivatized distearoyl phosphatidylethanolamine (PEG-DSPE). Methods of  
10 preparing PEG-PE are well known and typically involve using an activated methoxy PEG (with only one reactive end) and PE. Thus PEG-succinimidyl succinate may be reacted in a basic organic solvent (Klibanov *et al.*, *FEBS Lett.*, 268: 235-237 (1990)). A particularly preferred method of PEG-PE preparation is based on reaction of the PEG with carbonyldiimidazole followed by addition of PE  
15 (*see*, Woodle *et al.* *Proc. Intern. Symp. Control. Rel. Bioact. Mater.*, 17: 77-78 (1990), Papahadjopoulos *et al.*, *Proc. Natl. Acad. Sci. USA*, 88: 11460-11464 (1991), Allen *et al.*, *Biochim. Biophys. Acta.*, 1066: 29-36 (1991), Woodle *et al.*, *Biochim. Biophys. Acta.*, 1105: 193-200 (1992), and Woodle *et al.*, *Period. Biol.*, 93: 349-352 (1991)). Similarly, cyanuric chloride activated PEG in a basic  
20 organic solvent is described by Blume *et al.* *Biochim. Biophys. Acta.*, 1029: 91-97 (1990) and U.S. Patent No. 5,213,804 which is herein incorporated by reference. A completely different approach is based on coupling the PEG with preformed liposomes utilizing tresyl chloride activated PEG which is then added to liposomes containing PE at high pH (Senior *et al.*, *Biochim. Biophys. Acta.*, 1-62: 77-82  
25 91991). Derivatized PEG is also commercially available. Thus, for example, PEG-PE is available from Avanti Polar lipids (Alabaster, Alabama) or Liposome Technology (Menlo Park, California, USA). One of skill in the art will recognize that many other linkages are available.

### C) Fab' Antibody Fragment

In a preferred embodiment, the liposomes of the present invention are conjugated to the Fab' region of an antibody which acts as a targeting moiety enabling the liposome to specifically bind a target cell bearing the target molecule (e.g., characteristic marker) to which the Fab' antibody fragment is directed. In general terms, the Fab' region of an antibody represents a monomer comprising the variable regions and the C<sub>H</sub>1 region of one arm of an antibody.

An "antibody" refers to a protein consisting of one or more polypeptides substantially encoded by immunoglobulin genes or fragments of immunoglobulin genes. The recognized immunoglobulin genes include the kappa, lambda, alpha, gamma, delta, epsilon and mu constant region genes, as well as the myriad immunoglobulin variable region genes. Light chains are classified as either kappa or lambda. Heavy chains are classified as gamma, mu, alpha, delta, or epsilon, which in turn define the immunoglobulin classes, IgG, IgM, IgA, IgD and IgE, respectively.

The basic immunoglobulin (antibody) structural unit is known to comprise a tetramer. Each tetramer is composed of two identical pairs of polypeptide chains, each pair having one "light" (about 25 kD) and one "heavy" chain (about 50-70 kD). The N-terminus of each chain defines a variable region of about 100 to 110 or more amino acids primarily responsible for antigen recognition. The terms variable light chain (V<sub>L</sub>) and variable heavy chain (V<sub>H</sub>) refer to these light and heavy chains respectively.

Antibodies may exist as intact immunoglobulins or as a number of well characterized fragments produced by digestion with various peptidases. In particular, pepsin digests an antibody below the disulfide linkages in the hinge region to produce F(ab')<sub>2</sub>, a dimer of Fab which itself is a light chain joined to V<sub>H</sub>-C<sub>H</sub>1 by a disulfide bond. The F(ab')<sub>2</sub> may be reduced under mild conditions to break the disulfide linkage in the hinge region thereby converting the F(ab')<sub>2</sub> dimer into an Fab' monomer. The Fab' monomer is essentially an Fab with part of the hinge region (see, *Fundamental Immunology*, W.E. Paul, ed., Raven Press, N.Y. (1993) for more antibody fragment terminology). While the Fab' domain is

defined in terms of the digestion of an intact antibody, one of skill will appreciate that such Fab' fragments may be synthesized *de novo* either chemically or by utilizing recombinant DNA methodology.

The Fab' regions used in the present invention may be derived from antibodies of animal (especially mouse or rat) or human origin or may be chimeric (Morrison et al., *Proc Natl. Acad. Sci. USA* 81, 6851-6855 (1984) both incorporated by reference herein) or humanized (Jones et al., *Nature* 321, 522-525 (1986), and published UK patent application No. 8707252, both incorporated by reference herein).

The Fab' region is selected to specifically bind to a molecule or marker characteristic of the surface of the cells to which it is desired to deliver the contents of the liposome. A molecule is characteristic of cell, tissue, or physiological state when that molecule is typically found in association with that cell type or alternatively in a multiplicity of cell types all expressing a particular physiological condition (*e.g.*, transformation). A specific characteristic marker is preferably found on the surfaces of cells of a particular tissue or cell type or on the surfaces of tissues or cells expressing a particular physiological condition and on no other tissue or cell type in the organism. One of skill will recognize however, that such a level of specificity of the marker is often not required. For example a characteristic cell surface marker will show sufficient tissue specificity if the only non-target tissues are not accessible to the liposome. Alternatively, effective specificity may be achieved by overexpression of the marker in the target tissue as compared to other tissues. This will result in preferential uptake by the target tissue leading to effective tissue specificity. Thus for example, many cancers are characterized by the overexpression of cell surface markers such as the HER2 (*c-erbB-2*, *neu*) proto-oncogene encoded receptor in the case of breast cancer.

One of skill will recognize that there are numerous cell surface markers that provide good characteristic markers for liposomes depending on the particular tissue it is desired to target. These cell surface markers include, but are not limited to carbohydrates, proteins, glycoproteins, MHC complexes, and

receptor proteins such as HER, CD4 and CD8 receptor proteins as well as other growth factor receptor proteins.

Growth factor receptors are particularly preferred characteristic cell surface markers. Growth factor receptors are cell surface receptors that specifically bind growth factors and thereby mediate a cellular response characteristic of the particular growth factor. The term "growth factor", as used herein, refers to a protein or polypeptide ligand that activates or stimulates cell division or differentiation or stimulates biological response like motility or secretion of proteins. Growth factors are well known to those of skill in the art and include, but are not limited to, platelet-derived growth factor (PDGF), epidermal growth factor (EGF), insulin-like growth factor (IGF), transforming growth factor  $\beta$  (TGF- $\beta$ ), fibroblast growth factors (FGF), interleukin 2 (IL2), nerve growth factor (NGF), interleukin 3 (IL3), interleukin 4 (IL4), interleukin 1 (IL1), interleukin 6 (IL6), interleukin 7 (IL7), granulocyte/macrophage colony-stimulating factor (GM-CSF), granulocyte colony-stimulating factor (G-CSF), macrophage colony-stimulating factor (M-CSF), erythropoietin and the like. One of skill in the art recognizes that the term growth factor as used herein generally includes cytokines and colony stimulating factors.

Particularly preferred markers are found in the HER family of growth factor receptors. More specifically HER1, HER2, HER3 and HER4 are more preferred with HER2 most preferred. The HER receptors comprise protein tyrosine kinases that themselves provide highly specific antibody targets. Thus, in one embodiment, the P185 tyrosine kinase of HER2 provides a most preferred target for the Fab' antibody domain of the utilized in the immunoliposomes of the present invention.

It will be appreciated that the characteristic marker need not be a naturally occurring marker, but rather may be introduced to the particular target cell. This may be accomplished by directly tagging a cell or tissue with a particular marker (e.g., by directly injecting the particular target tissue with a marker, or alternatively, by administering to the entire organism a marker that is selectively incorporated by the target tissue. In one embodiment, the marker may

be a gene product that is encoded by a nucleic acid in an expression cassette. The marker gene may be under the control of a promoter that is active only in the particular target cells. Thus introduction of a vector containing the expression cassette will result in expression of the marker in only the particular target cells.

- 5 One of skill in the art will recognize that there are numerous approaches utilizing recombinant DNA methodology to introduce characteristic markers into target cells.

In one preferred embodiment, the targeting moiety will specifically bind products or components of a growth factor receptor, in particular products of  
10 the HER2 (*c-erbB-2*, *neu*) proto-oncogene. It is particularly preferred that the targeting moiety bind the growth factor receptor-tyrosine kinase encoded by HER2, protein p185<sup>HER2</sup>, which is commonly overexpressed in breast cancers (Slamon *et al.*, *Science*, 235: 177-182 (1987). Other suitable targets for the targeting moiety include, but are not limited to EGFR (HER1), HER3, and HER4, combinations of  
15 these receptors, and other markers associated with cancers. Other antibodies of interest include, but are not limited to BR96 (Friedman *et al.*, *Cancer Res.*, 53: 334-339 (1993), e23 to erbB2 (Batra *et al.*, *Proc. Natl. Acad. Sci. USA*, 89: 5867-5871 (1992)), PR1 in prostate cancer (Brinkmann *et al.*, *Proc. Natl. Acad. Sci. USA.*, 90: 547-551 (1993)), and K1 in ovarian cancer (Chang *et al.* *Int. J. Cancer*,  
20 50: 373-381 (1992).

Immunoliposomes of the present invention may be prepared by incorporating the Fab' antibody domain moieties into the liposomes by a variety of techniques well known to those of skill in the art. For example, a biotin conjugated Fab' may be bound to a liposome containing a streptavidin.  
25 Alternatively, the biotinylated Fab' may be conjugated to a biotin derivatized liposome by an avidin or streptavidin linker. Thus, for example, a biotinylated monoclonal antibody was biotinylated and attached to liposomes containing biotinylated phosphatidylethanolamine by means of an avidin linker. (See, for example, Ahmad *et al.*, *Cancer Res.*, 52: 4817-4820 (1992) which is herein  
30 incorporated by reference). Typically about 30 to 125 and more typically about 50 to 100 Fab' molecules per liposome are used.

In a preferred embodiment, the targeting moiety may be directly conjugated to the liposome. Such means of direct conjugation are well known to those of skill in the art. See for example, G. Gregoriadis, (1984) "Liposome Technology" CRC Press, Boca Raton, Florida and D.D. Lasic, "Liposomes: from  
5 physics to applications" (1993) Elsevier, Amsterdam; N.Y. Particularly preferred is conjugation through a thioether linkage. This may be accomplished by reacting the antibody with a maleimide derivatized lipid such as maleimide derivatized phosphatidylethanolamine (M-PE) or dipalmitoylethanolamine (M-DEP). This approach is described in detail by Martin *et al.* *J. Biol. Chem.*, 257: 286-288  
10 (1982) which is incorporated herein by reference.

#### Preparation of Liposomes

A variety of methods are available for preparing liposomes as described in, *e.g.*, Szoka *et al.*, *Ann. Rev. Biophys. Bioeng.* 9:467 (1980), U.S.  
15 Pat. Nos. 4,186,183, 4,217,344, 4,235,871, 4,261,975, 4,485,054, 4,501,728, 4,774,085, 4,837,028, 4,946,787, PCT Publication No. WO 91/17424, Szoka & Papahadjopoulos, *Proc. Natl. Acad. Sci. USA*, 75: 4194-4198 (1978), Deamer and Bangham, *Biochim. Biophys. Acta*, 443: 629-634 (1976); Fraley, *et al.*, *Proc. Natl. Acad. Sci. USA*, 76: 3348-3352 (1979); Hope, *et al.*, *Biochim. Biophys. Acta*, 812:  
20 55-65 (1985); Mayer, *et al.*, *Biochim. Biophys. Acta*, 858: 161-168 (1986); Williams, *et al.*, *Proc. Natl. Acad. Sci.*, 85: 242-246 (1988), the text *Liposomes*, Marc J. Ostro, ed., Marcel Dekker, Inc., New York, 1983, Chapter 1, and Hope, *et al.*, *Chem. Phys. Lip.* 40: 89 (1986), all of which are incorporated herein by reference. Suitable methods include, *e.g.*, sonication, extrusion, high  
25 pressure/homogenization, microfluidization, detergent dialysis, calcium-induced fusion of small liposome vesicles, and ether-infusion methods, all well known in the art. One method produces multilamellar vesicles of heterogeneous sizes. In this method, the vesicle-forming lipids are dissolved in a suitable organic solvent or solvent system and dried under vacuum or an inert gas to form a thin lipid film.  
30 If desired, the film may be redissolved in a suitable solvent, such as tertiary butanol, and then lyophilized to form a more homogeneous lipid mixture which is



in a more easily hydrated powder-like form. This film is covered with an aqueous buffered solution and allowed to hydrate, typically over a 15-60 minute period with agitation. The size distribution of the resulting multilamellar vesicles can be shifted toward smaller sizes by hydrating the lipids under more vigorous agitation  
5 conditions or by adding solubilizing detergents such as deoxycholate.

In a preferred embodiment, multilamellar liposomes are produced by the reverse phase evaporation method of Szoka & Papahadjopoulos, *Proc. Natl. Acad. Sci. USA*, 75: 4194-4198 (1978).

Unilamellar vesicles are generally prepared by sonication or  
10 extrusion. Sonication is generally performed with a tip sonifier, such as a Branson tip sonifier, in an ice bath. Typically, the suspension is subjected to several sonication cycles. Extrusion may be carried out by biomembrane extruders, such as the Lipex Biomembrane Extruder. Defined pore size in the extrusion filters may generate unilamellar liposomal vesicles of specific sizes. The liposomes may  
15 also be formed by extrusion through an asymmetric ceramic filter, such as a Ceraflow Microfilter, commercially available from the Norton Company, Worcester MA.

Following liposome preparation, the liposomes which have not been sized during formation may be sized to achieve a desired size range and relatively  
20 narrow distribution of liposome sizes. A size range of about 0.2-0.4 microns allows the liposome suspension to be sterilized by filtration through a conventional filter, typically a 0.22 micron filter. The filter sterilization method can be carried out on a high through-put basis if the liposomes have been sized down to about 0.2-0.4 microns.

Several techniques are available for sizing liposomes to a desired  
25 size. One sizing method is described in U.S. Pat. Nos. 4,529,561 or 4,737,323, incorporated herein by reference. Sonicating a liposome suspension either by bath or probe sonication produces a progressive size reduction down to small unilamellar vesicles less than about 0.05 microns in size. Homogenization is  
30 another method which relies on shearing energy to fragment large liposomes into smaller ones. In a typical homogenization procedure, multilamellar vesicles are

recirculated through a standard emulsion homogenizer until selected liposome sizes, typically between about 0.1 and 0.5 microns, are observed. The size of the liposomal vesicles may be determined by quasi-electric light scattering (QELS) as described in Bloomfield, *Ann. Rev. Biophys. Bioeng.*, 10: 421-450 (1981),  
5 incorporated herein by reference. Average liposome diameter may be reduced by sonication of formed liposomes. Intermittent sonication cycles may be alternated with QELS assessment to guide efficient liposome synthesis.

Extrusion of liposome through a small-pore polycarbonate membrane or an asymmetric ceramic membrane is also an effective method for reducing  
10 liposome sizes to a relatively well-defined size distribution. Typically, the suspension is cycled through the membrane one or more times until the desired liposome size distribution is achieved. The liposomes may be extruded through successively smaller-pore membranes, to achieve a gradual reduction in liposome size. For use in the present invention, liposomes having a size of about 0.05  
15 microns to about 0.15 microns. More preferred are liposomes having a size of about 0.05 to 0.5 microns.

### Immunoliposome Contents

The therapeutic agent which may be used is any compound including  
20 the ones listed below which can be stably entrapped in liposomes at a suitable loading factor and administered at a therapeutically effective doses (indicated below in parentheses after each compound,  $m^2$  refers to body surface area). These include amphipathic antitumor compounds such as the plant alkaloids vincristine ( $1.4 \text{ mg}/m^2$ ), vinblastine ( $4-18 \text{ mg}/m^2$ ) and etoposide ( $35-100 \text{ mg}/m^2$ ), and the  
25 anthracycline antibiotics including doxorubicin ( $60-75 \text{ mg}/m^2$ ), epirubicin ( $60-120 \text{ mg}/m^2$ ) and daunorubicin ( $25-45 \text{ mg}/m^2$ ). The water-soluble anti-metabolites such as methotrexate ( $3 \text{ mg}/m^2$ ), cytosine arabinoside ( $100 \text{ mg}/m^2$ ) and fluorouracil ( $10-15 \text{ mg}/\text{kg}$ ), the antibiotics such as bleomycin ( $10-20 \text{ units}/m^2$ ), mitomycin ( $20 \text{ mg}/m^2$ ), plicamycin ( $25-30 \mu\text{g}/m^2$ ) and dactinomycin ( $15 \mu\text{g}/m^2$ ), and the alkylating  
30 agents including cyclophosphamides and derivatives thereof ( $3-25 \text{ mg}/\text{kg}$ ), thiotepea ( $0.3-0.4 \text{ mg}/\text{kg}$ ) and BCNU ( $150-200 \text{ mg}/m^2$ ) are also useful in this context.

Other suitable drugs include aclacinomycin, idarubicin, mitoxantrone, cisplatin and other Platinum II analogs. The liposomes may also contain the taxanes including taxol, taxotere, dihydroxytaxanes, camptothecines and other taxane derivatives and isolates. In addition, the liposomes may contain encapsulated tumor-therapeutic peptides (e.g., plant or bacterially derived toxins) and protein drugs such as IL-2 and/or TNF, and/or immunomodulators, such as M-CSF, which are present alone or in combination with anti-tumor drugs, such as anthracycline antibiotic drugs. The immunoliposomes may contain fluorinated pyrimidine and purine bases or nucleosides. The immunoliposomes may also contain nucleic acids such as oligonucleotides containing natural or modified bases and having a phosphodiester internucleotide linkage or modified internucleotide linkages such as a phosphorothioate or polyamide linkages. One of skill will recognize that nucleic acids may be used as antisense or triplex-forming molecules to block transcription and translation through binding of DNA or RNA. Alternatively, the nucleic acids may be used to transform cells and to induce the expression of heterologous proteins. In this latter context, the nucleic acid will comprise an expression cassette which includes the nucleic acid sequence encoding the protein to be expressed under the control of a promoter.

#### 20 Loading Therapeutic Compositions into Immunoliposomes

The methods of loading conventional drugs into liposomes are well known to those of skill in the art. The most common methods include an encapsulation technique and the transmembrane potential loading method. In the encapsulation technique, the drug is placed into the buffer from which the liposomes are made. The latter method has been described in detail in U.S. Patent No. 4,885,172, U.S. Patent No. 5,059,421, and U.S. Patent No. 5,171,578, the contents of which are incorporated herein by reference.

Briefly, the transmembrane potential loading method can be used with essentially any conventional drug which can exist in a charged state when dissolved in an appropriate aqueous medium. Preferably, the drug will be relatively lipophilic so that it will partition into the liposome membranes. A

transmembrane potential is created across the bilayers of the liposomes or targeting moiety liposome conjugates and the drug is loaded into the liposome by means of the transmembrane potential. The transmembrane potential is generated by creating a concentration gradient for one or more charged species (*e.g.*, Na<sup>+</sup>, K<sup>+</sup> and/or H<sup>+</sup>) across the membranes. This concentration gradient is generated by producing liposomes or targeting moiety liposome conjugates having different internal and external media. Thus, for a drug which is positively charged when ionized, a transmembrane potential is created across the membranes which has an inside potential which is negative relative to the outside potential, while for a drug which is negatively charged, the opposite orientation is used.

#### Assaying Blood Half-life.

One of the requirements for liposome localization in a target tissue is an extended immunoliposome lifetime in the bloodstream following administration. One measure of immunoliposome lifetime in the bloodstream is the blood/RES ratio determined at a selected time after liposome administration. Typically immunoliposomes containing a label (*e.g.* fluorescent marker, electron dense reagent, or radioactive marker), either internal in the liposome or bound to a lipid comprising the liposome, are injected into the test organism. A fixed period of time later, the organism is sacrificed and the amount of label detected in the blood (*e.g.* by measuring luminescence, or scintillation counting) is compared to that localized in particular tissues (*e.g.* liver or spleen).

The time course of retention of immunoliposomes in the blood may also simply be determined by sampling blood at fixed intervals after administration of label-containing liposomes and determining the amount of label remaining in the circulation. The result may be expressed as the fraction of the original dose.

#### Assaying Uptake Into the Cytoplasm of Target Cells and Determining Tissue Distribution

Uptake and internalization of immunoliposomes into the cytoplasm of target cells may similarly be determined by administering immunoliposomes

containing a label (*e.g.* fluorescent marker, electron dense reagent, or radioactive marker) and subsequently detecting the presence or absence of that label in the cytoplasm of the target cell. For example an immunoliposome containing a fluorescent marker, such as rhodamine conjugated to the lipid constituting the liposome itself, may be administered to the organism or simply to cells in culture. The tissues or cells may then be fixed and the fluorescence detected using fluorescence microscopy. Similarly, an electron-dense label (*e.g.* gold) may be used and detected using electron microscopy. One of skill in the art will recognize that many labels are suitable and the method of detection will reflect the choice of label.

#### Assaying for Antiproliferative Activity of Immunoliposomes

The present invention provides for growth-inhibiting immunoliposomes that essentially comprise an empty immunoliposome having an Fab' targeting moiety directed to a cell surface receptor. Particularly preferred are growth factor receptors. Identification of Fab' immunoliposomes that are particularly effective inhibitors of cell proliferation may be accomplished with routine screening. This involves providing a cell culture where the cells bear a growth factor receptor, or other characteristic cell surface marker, to which the Fab' fragment is directed, contacting the cells in the culture with the immunoliposome to be tested, and measuring the resulting change in cell proliferation rate. Means of measuring cell proliferation rate are well known to those of skill in the art.

In one approach, for example, proliferation rate may be assessed directly by measuring the change in actual numbers of cells over a fixed period of time. Thus, as illustrated in Example 3, tumor cells such as SK-BR-3 or BT-474 cells were grown in monolayer culture and then incubated at 37°C with varying concentrations of immunoliposomes based on antibody content. After continuous treatment for 4 days, cell monolayers were washed with PBS and stained with crystal violet dye (0.5% in methanol) for determination of relative proliferation as

previously described (Hudziak, *et al.*, *Mol. Cell Biol.* 9: 1165-1172 (1989) which is incorporated herein by reference).

Alternatively, it is well known that increases in cell proliferation rate are typically accompanied by increases in metabolic rate. Thus proliferation may be evaluated indirectly by measuring changes in metabolic rate of cells exposed to the immunoliposome to be tested. Numerous means of measuring metabolic rate are well known to those of skill in the art. On particularly preferred approach is to measure the rate of uptake of a labeled metabolic precursor such as tritiated thymidine. Briefly this is accomplished by administering the [<sup>3</sup>H]-thymidine to a test culture containing the immunoliposome and to a control culture lacking the immunoliposome. After a fixed period of time, cells are collected and the amount of [<sup>3</sup>H]-thymidine taken up by the cells is measured utilizing standard techniques (e.g., scintillation counting). Comparison of the test and control cells indicates changes in metabolic activity and therefore proliferation rate.

#### Pharmaceutical Compositions

Pharmaceutical compositions comprising the immunoliposomes of the invention are prepared according to standard techniques and further comprise a pharmaceutically acceptable carrier. Generally, normal saline will be employed as the pharmaceutically acceptable carrier. Other suitable carriers include, e.g., water, buffered water, 0.4% saline, 0.3% glycine, and the like, including glycoproteins for enhanced stability, such as albumin, lipoprotein, globulin, etc. These compositions may be sterilized by conventional, well known sterilization techniques. The resulting aqueous solutions may be packaged for use or filtered under aseptic conditions and lyophilized, the lyophilized preparation being combined with a sterile aqueous solution prior to administration. The compositions may contain pharmaceutically acceptable auxiliary substances as required to approximate physiological conditions, such as pH adjusting and buffering agents, tonicity adjusting agents and the like, for example, sodium acetate, sodium lactate, sodium chloride, potassium chloride, calcium chloride, *etc.* Additionally, the liposome suspension may include lipid-protective agents which protect lipids

against free-radical and lipid-peroxidative damages on storage. Lipophilic free-radical quenchers, such as alphotocopherol and water-soluble iron-specific chelators, such as ferrioxamine, are suitable.

The concentration of immunoliposomes, in the pharmaceutical formulations can vary widely, *i.e.*, from less than about 0.05%, usually at or at least about 2-5% to as much as 10 to 30% by weight and will be selected primarily by fluid volumes, viscosities, *etc.*, in accordance with the particular mode of administration selected. For xample, the concentration may be increased to lower the fluid load associated with treatment. This may be particularly desirable in patients having atherosclerosis-associated congestive heart failure or severe hypertension. Alternatively, immunoliposomes composed of irritating lipids may be diluted to low concentrations to lessen inflammation at the site of administration. The amount of immunoliposome administered will depend upon the particular Fab' used, the disease state being treated, the therapeutic agent being delivered, and the judgement of the clinician. Generally the amount of immunoliposomes administered will be sufficient to deliver a therapeutically effective dose of the particular pharmacological agent. The quantity of immunoliposomes necessary to deliver a therapeutically effective dose can be determined by uptake assays as described above. Therapeutically effective dosages for various pharmacological agents are well known to those of skill in the art and representative ranges are given for a number of pharmaceuticals above. Typical immunoliposome dosages will generally be between about 0.01 and about 50 mg per kilogram of body weight, preferably between about 0.1 and about 10 mg/kg of body weight.

Preferably, the pharmaceutical compositions are administered parenterally, *i.e.*, intraarticularly, intravenously, intraperitoneally, subcutaneously, or intramuscularly. More preferably, the pharmaceutical compositions are administered intravenously or intraperitoneally by a bolus injection. Particular formulations which are suitable for this use are found in *Remington's Pharmaceutical Sciences*, Mack Publishing Company, Philadelphia, PA, 17th ed. (1985). Typically, the formulations will comprise a solution of the liposomes

suspended in an acceptable carrier, preferably an aqueous carrier. A variety of aqueous carriers may be used, *e.g.*, water, buffered water, 0.9% isotonic saline, and the like. These compositions may be sterilized by conventional, well known sterilization techniques, or may be sterile filtered. The resulting aqueous solutions  
5 may be packaged for use as is, or lyophilized, the lyophilized preparation being combined with a sterile aqueous solution prior to administration. The compositions may contain pharmaceutically acceptable auxiliary substances as required to approximate physiological conditions, such as pH adjusting and buffering agents, tonicity adjusting agents, wetting agents and the like, for example, sodium acetate,  
10 sodium lactate, sodium chloride, potassium chloride, calcium chloride, sorbitan monolaurate, triethanolamine oleate, etc.

### EXAMPLES

The invention is illustrated by the following examples. These  
15 examples are offered to illustrate, but not to limit the present invention .

#### Example 1

##### Preparation of Liposomes and Immunoliposomes

##### A) Materials

20 Egg phosphatidylcholine (EPC) was purchased from Avanti Polar Lipids (Alabaster, AL); cholesterol (Chol) from Calbiochem (San Diego, CA); N-Tris[hydroxymethyl]-2-aminoethanesulfonic acid (TES) from Sigma; hydrogenated soy phosphatidylcholine (HSPC) from Natterman (Cologne, FRG); rhodamine-labeled phospholipids from Avanti; desferrioxamine mesylate (desferal)  
25 from Ciba-Geigy (Summit, NJ); doxorubicin from Farmitalia, Carlo Erba (Milan, Italy) or Cetus (Emeryville, CA); and N-[4*p*-maleimidophenyl]butyryl]phosphatidylethanolamine (M-PE) from Molecular Probes (Portland, OR). PEG ( $M_r = 1900$ )-derivatized phosphatidylethanolamine (PEG-PE) was synthesized as described (Allen *et al.*, *Biochim. Biophys. Acta*,  
30 1066: 29-36 (1991)), and obtained from Liposome Technology, Inc. (Menlo Park, CA).



### B) Preparation of Fab' Fragments.

Cloned rhuMAbHER2 sequences for heavy and light chain were co-expressed in *E. coli* as previously described (Carter *et al.*, *Biotechnology* 10: 163-167 (1992)). The antibody fragment, rhuMAbHER2-Fab', was recovered  
5 from *E. coli* fermentation pastes by affinity chromatography with Streptococcal protein G (Carter *et al.*, *Biotechnology*, 10:163-167 (1992) which is incorporated herein by reference), typically yielding Fab' with 60-90% containing reduced free thiol (Fab'-SH). As a control, an irrelevant humanized Fab' was used. Derived from an anti-CD 18 murine monoclonal antibody, rhuMAbH52-Fab' differs from  
10 rhuMAbHER2-Fab' only by replacement of the antigen-binding loops, and showed no detectable binding to any known murine or human antigen (Eigenbrot *et al. Proteins: Structure, Function, and Genetics*, 18: 49-62 (1994)).

### C) Preparation of Liposomes.

Liposomes were prepared according to the reverse phase evaporation  
15 method (Szoka & Papahadjopoulos, *Proc. Natl. Acad. Sci. USA*, 75: 4194-4198 (1978), with lipid composition including EPC:Chol (2:1) or, where stated, HSPC:Chol (3:2) and PEG-PE (0-6 mole%). Liposomes were subsequently extruded repeatedly under positive pressure with argon gas through polycarbonate membrane filters of defined pore size sequentially from 0.1 to 0.05  $\mu$ m (Olson, *et al.*, *Biochim. Biophys. Acta*, 55: 9-23 (1979); Szoka *et al.*, *Biochem. Biophys. Acta*, 601: 559-571 (1980)). This procedure yields liposomes of 60-120 nm diameter as determined by dynamic light scattering. Liposome concentration was determined by phosphate assay (Bartlett, *J. Biol. Chem.*, 234: 466-468 (1959)). For immunoliposome preparations, 2 mole% M-PE (of total phospholipid) was  
25 included in the lipid mixture in chloroform prior to the making of liposomes (Martin and Papahadjopoulos, *J. Biol. Chem.*, 257: 286-288 (1982)). Liposomes without encapsulated doxorubicin were prepared in HEPES-NaCl buffer, pH 7.2, 300 mOsm. HSPC/Chol liposomes containing doxorubicin were prepared in 250 mM ammonium sulfate containing 1 mM desferal at pH 5.5. Unencapsulated  
30 ammonium sulfate was removed by gel filtration with G-75 Sephadex. Doxorubicin in powder form was then dissolved in this liposome suspension at 0.1

mg doxorubicin/ $\mu$ mole phospholipid(Papahadjopoulos *et al.*, *Proc. Natl. Acad. Sci. USA*, 88: 11460-11464 (1991); Huang *et al.*, *Cancer Res.*, 52:6774-6781 (1992)), which traps doxorubicin within the interior space of the liposome (Lasic *et al.*, *FEBS Lett.*, 312: 255-258 (1992)). The efficiency of drug loading via salt gradient  
5 was high and reached >99% loading when 1 mg drug per 10  $\mu$ mole phospholipid was used.

Fab' was conjugated to the liposomes after drug loading via thioether linkage, as previously described(Martin *et al. J. Biol. Chem.*, 257: 286-288 (1982). Since maleimide is more stable at lower pH, all procedures were performed at pH  
10 5.5. Unreacted Fab' was separated from immunoliposomes by gel filtration with Sephacryl S-400. The maleimide group on immunoliposomes was deactivated after conjugation by 2-fold excess of mercaptoethanol to M-PE. The amount of Fab' conjugated was determined by BioRad protein assay.

Four types of liposomes were prepared. "Conventional" liposomes  
15 without antibody were composed of phosphatidylcholine and cholesterol only. "Sterically stabilized" liposomes additionally contained PEG-PE. Immunoliposomes were prepared by conjugation of the above with Fab' fragments derived from the humanized antibody rhuMAbHER2 to yield conventional or sterically stabilized immunoliposomes. Fab' fragments rather than intact antibody  
20 were used for the following reasons: 1) rhuMAbHER2-Fab' fragments can be expressed as recombinant proteins in *E. coli* at extremely high efficiency (Carter *et al.*, *supra.* ) the free thiol group in the Fab' hinge region provides a readily available site for covalent attachment to modified liposomes, and is distant from the antigen binding site; and 3) rhuMAbHER2-Fab' has far less antiproliferative  
25 activity than intact rhuMAbHER2, and thus it was of interest to see whether attachment of anti-p185<sup>HER2</sup>-Fab' to liposomes would reconstitute this activity. Typically, conjugation resulted in approximately 50-100 Fab' molecules per liposome particle.

## Example 2

### Immunoliposome Binding

The ability of anti-p185<sup>HER2</sup> immunoliposomes to bind *in vitro* to breast cancer cells which overexpress the p185<sup>HER2</sup> receptor was evaluated by two methods; a flow cytometric assay and a competitive binding assay. For the flow cytometric assay, SK-BR-3, which express high levels of p185<sup>HER2</sup>, or MCF-7 cells, were exposed to anti-p185<sup>HER2</sup> immunoliposomes for 45 minutes on ice, washed with PBS, stained with a secondary anti-human antibody to detect bound immunoliposomes (FITC-labeled goat anti-human IgG), washed with PBS again, and then subjected to flow cytometry (Fig. 1). SK-BR-3 cells bound significant amounts of either conventional or sterically stabilized anti-p185<sup>HER2</sup> immunoliposomes, but not control liposomes lacking Fab'. MCF-7 breast cancer cells, which do not overexpress p185<sup>HER2</sup>, showed minimal binding to anti-p185<sup>HER2</sup> immunoliposomes (data not shown).

Another measure of binding was afforded by a competitive binding assay, in which SK-BR-3 (breast cancer cells) or BT-474 cells in monolayer culture were simultaneously incubated with <sup>125</sup>I-labeled rhuMabHER2 or muMab4D5, at 0.1 nM for 18 hours at 4°C, and increasing concentrations of anti-p185<sup>HER2</sup> immunoliposomes (Fig. 2). Counts bound were determined by gamma counting. Anti-p185<sup>HER2</sup> immunoliposomes efficiently displaced binding of rhuMabHER2 to both SK-BR-3 cells (data not shown) and BT-474 cells (which also express high levels of p185<sup>HER2</sup>). An approximation of binding affinity was obtained via Scatchard analysis of the binding data, by making the assumption that Fab' on immunoliposomes behaved as free ligand. Using this model, apparent binding constants for the immunoliposomes were comparable to those of free (i.e. non-liposomal) rhuMabHER2-Fab' or intact rhuMabHER2. Conventional or sterically stabilized (6 mole% PEG-PE) control liposomes lacking Fab' showed negligible binding. A summary of the binding data is shown in Table 1.

Table 1. Binding of anti-P185<sup>HER2</sup> immunoliposomes to BT-474 cells.

5	Ligand	Apparent $K_d$ (nM)	$EC_{50}$ (nM)
	Control Liposome, Conventional	No displacement	>1000
	Control Liposome, Sterically Stabilized	2500	>1000
	Anti-p185 <sup>HER2</sup> Immunoliposome, Conventional	9	15
10	anti-p185 <sup>HER2</sup> Immunoliposome, Sterically Stabilized	58	78
	rhuMabHER2-Fab'	22	30
	rhuMabHER2	1	2

15

**Example 3****Anti-Proliferative Activity of "Growth-Inhibiting" Immunoliposomes**

To test the antiproliferative activity of immunoliposomes alone, without encapsulated drugs, tumor cells such as SK-BR-3 or BT-474 cells were grown in monolayer culture and then incubated at 37°C with varying concentrations of immunoliposomes based on antibody content. After continuous treatment for 4 days, cell monolayers were washed with PBS and stained with crystal violet dye (0.5% in methanol) for determination of relative proliferation as previously described (Hudziak, *et al.*, *Mol. Cell Biol.* 9: 1165-1172 (1989)).

While intact (bivalent) rhuMabHER2 inhibited the growth of p185<sup>HER2</sup>-overexpressing breast cancer cells in monolayer culture, monovalent Fab' fragments of this antibody (rhuMabHER2-Fab') are much less effective at inhibiting growth (O'Connell *et al.* pages 218-239 In *Protein Folding In Vivo and In Vitro.*, Cleland JL, ed. Washington, D.C., American Chemical Society, (1993)). This observation suggested that cross-linking of p185<sup>HER2</sup> receptors by bivalent antibody is important for the antiproliferative effect, and raised the question of whether liposomal anchoring of rhuMabHER2-Fab' fragments might improve the antiproliferative activity of Fab' by increasing effective valency.

The effect of anti-p185<sup>HER2</sup> immunoliposomes upon SK-BR-3 cells in monolayer culture was tested and compared to rhuMabHER2 and

5 rhuMabHER2-Fab' (Fig.3). Treatment with conventional or sterically stabilized control liposomes lacking Fab' did not significantly affect cell growth. In contrast, both conventional and sterically stabilized anti-p185<sup>HER2</sup> immunoliposomes inhibited growth in a dose-dependent manner. The growth inhibitory effect of the immunoliposomes reached a plateau of about 30% growth inhibition (70% of control growth), approaching the 40% growth inhibition seen with free intact rhuMabHER2. In contrast, free rhuMabHER2-Fab' induced only modest growth inhibition. Similar results were also obtained with p185<sup>HER2</sup>-overexpressing BT-474 breast cancer cells (data not shown).

10 It is notable that liposome-associated rhuMabHER2-Fab' produced a markedly greater antiproliferative effect than the same amount of rhuMabHER2-Fab' free in solution. A plausible explanation is that liposomal anchoring of rhuMabHER2-Fab' allows cross-linking of p185<sup>HER2</sup>, thus yielding biological activity which is comparable to that of the intact bivalent antibody.

15

#### Example 4

##### Cytotoxicity of Anti-p185<sup>HER2</sup> Immunoliposomes Containing Doxorubicin.

Although empty anti-p185<sup>HER2</sup> immunoliposomes displayed antiproliferative activity against p185<sup>HER2</sup>-overexpressing breast cancer cells in culture, it was possible to greatly augment the antineoplastic effect of the immunoliposomes by loading them with cytotoxic agents, thus producing a targeted drug delivery system. Doxorubicin was used, because of preclinical and clinical evidence suggesting that doxorubicin may be particularly useful against breast cancers overexpressing p185<sup>HER2</sup>, with or without concomitant immunotherapy. It was therefore of interest to evaluate the cytotoxicity and specificity of doxorubicin-loaded anti-p185<sup>HER2</sup> immunoliposomes against p185<sup>HER2</sup>-overexpressing breast cancer cells and against non-malignant cells that do not overexpress p185<sup>HER2</sup>.

30 Because of the efficient internalization shown by p185<sup>HER2</sup> immunoliposomes (see Example 5), it was expected that the liposomes might be just as effective at killing p185<sup>HER2</sup>-overexpressing breast cancer cells in culture as

free doxorubicin, a small (MW 544) amphipathic molecule that readily passes through the cell membrane. On the other hand, cells which do not overexpress p185<sup>HER2</sup>, while susceptible to free doxorubicin, would escape cytotoxic injury from doxorubicin-loaded anti-p185<sup>HER2</sup> immunoliposomes because of the inability of the  
5 immunoliposomes to target them.

To test the cytotoxicity of doxorubicin-loaded immunoliposomes, SK-BR-3 or WI-38 cells in monolayer culture were incubated with free doxorubicin or doxorubicin-loaded immunoliposomes for 1 hour, and then washed extensively with media. The cells were then further incubated at 37°C for 3 days, after which  
10 cell number was estimated by crystal violet staining as described above. Comparison with other assays of cell growth including alamar blue staining, MTT staining, and direct cell counting yielded essentially the same results.

The results of the SK-BR-3 cells treated for 1 hour with various doxorubicin-loaded immunoliposome preparations, are shown in Figure 4. Under  
15 these conditions, the antiproliferative effect of rhuMAbHER2 is not apparent, as it requires continuous exposure of the cells to rhuMAbHER2 (Hudziak *et al.*, *Mol. Cell Biol.*, 9: 1165-1172 (1989)). Treatment with free doxorubicin for 1 hour yielded significant cytotoxicity, with an IC<sub>50</sub> of approximately 0.3 µg/ml. Doxorubicin-loaded anti-p185<sup>HER2</sup> immunoliposomes showed comparable  
20 dose-dependent cytotoxicity, with an IC<sub>50</sub> of approximately 0.2 µg/ml for conventional anti-p185<sup>HER2</sup> immunoliposomes and approximately 1.0 µg/ml for sterically stabilized (2 mole% PEG-PE) anti-p185<sup>HER2</sup> immunoliposomes. These results indicated that anti-p185<sup>HER2</sup> immunoliposome delivery of doxorubicin to p185<sup>HER2</sup>-overexpressing cells in culture was as efficient a process as the rapid  
25 diffusion of free doxorubicin into the cells. Doxorubicin-loaded anti-p185<sup>HER2</sup> immunoliposomes were between 10- to 30-fold more cytotoxic than doxorubicin-loaded immunoliposomes bearing irrelevant Fab', which only affected cell growth at relatively high concentrations (>3.3 µg/ml).

WI-38 cells, a non-malignant lung fibroblast cell line which  
30 expresses minimal levels of p185<sup>HER2</sup>, were also treated with doxorubicin and with doxorubicin-loaded anti-p185<sup>HER2</sup> immunoliposomes (Figure 4B). Free doxorubicin

again produced significant dose-dependent cytotoxicity against WI-38 cells. However, doxorubicin-loaded anti-p185<sup>HER2</sup> immunoliposomes produced much reduced (20-fold less) cytotoxicity against these cells, and were indistinguishable from doxorubicin-loaded immunoliposomes bearing irrelevant Fab'. These results  
5 demonstrated the specificity of anti-p185<sup>HER2</sup> immunoliposome treatment for p185<sup>HER2</sup>-overexpressing targets, and further confirmed that the cytotoxicity observed against SK-BR-3 cells was not simply due to leakage of doxorubicin out of anti-p185<sup>HER2</sup> immunoliposomes and into solution.

### 10 Example 5

#### Internalization of Immunoliposomes Into the Cytoplasm of the Target Cell

To assess internalization of immunoliposomes into the cytoplasm of the target cell by fluorescence microscopic assay, liposomes and immunoliposomes were prepared as described in Example 1 with the addition of rhodamine-phosphatidylethanolamine at 1 mole% of the phospholipid components. The  
15 resulting rhodamine-labeled liposomes or immunoliposomes were incubated for varying amounts of time at 37°C with SK-BR-3 cells grown to subconfluence on cover slips. The cells were then fixed with 3% paraformaldehyde, mounted in 90% glycerol/100 mM Tris, pH8.5, containing 0.1% p-phenylenediamine (Sigma)  
20 as an antibleaching reagent, and observed with a Leitz Aristoplan fluorescence microscope or a Molecular Dynamics MultiProbe 2001 confocal microscope.

To assess internalization and intracellular disposition by electron microscopy, immunoliposomes were loaded with colloidal gold particles of 5-15 nm as previously described(Huang *et al.*, *Cancer Res.*, 52: 5135-5143 (1992);  
25 Straubinger *et al.*, *Cell*, 32: 10639-1079 (1983). Gold-containing immunoliposomes were incubated at 37°C with SK-BR-3 cells grown on cover slips for varying amounts of time, and the cells were then fixed and processed for electron microscopy. Stabilization of liposomes was achieved using tannic acid in the primary fixation(Straubinger *et al.*, *supra*), which provided adequate albeit not  
30 optimal preservation of the ultrastructure.

The antibody rhuMAbHER2 is rapidly internalized by p185<sup>HER2</sup>-overexpressing tumor cells via receptor-mediated endocytosis (Sarup *et al.*, *Growth Regul.* 1: 72-82 (1991)). To assess whether anti-p185<sup>HER2</sup> immunoliposomes are internalized within SK-BR-3 cells, cells were treated with rhodamine-labeled immunoliposomes for different time intervals, fixed and visualized by fluorescence microscopy. SK-BR-3 cells treated with conventional or sterically stabilized control liposomes lacking Fab' showed neither surface nor internal rhodamine fluorescence, consistent with the inability of control liposomes to bind to these cells. When treated with conventional anti-p185<sup>HER2</sup> immunoliposomes, SK-BR-3 cells exhibited intense foci of fluorescence both at the cell surface and intracellularly within 30 minutes of treatment. Confocal fluorescence microscopy confirmed that rhodamine fluorescence was present both at the cell surface and internalized within the cytoplasm of SK-BR-3 cells. In contrast, treatment with sterically stabilized anti-p185<sup>HER2</sup> immunoliposomes containing high PEG-PE concentrations (6 mole%) resulted in minimal intracellular fluorescence after 30 minutes. Because it appeared that the presence of PEG-PE retarded immunoliposome internalization, sterically stabilized immunoliposomes containing reduced concentrations of PEG-PE were evaluated. Immunoliposomes containing 2 mole% PEG-PE yielded an intermediate degree of intracellular fluorescence after 30 minutes, *i.e.* less than that seen with conventional immunoliposomes but more than that seen with 6 mole% PEG-PE-containing immunoliposomes. Despite somewhat retarded internalization, the sterically stabilized immunoliposomes containing 2% PEG-PE did accumulate intracellularly with longer incubation time, such as at 2 hours. Thus, while anti-p185<sup>HER2</sup> immunoliposomes were internalized within SK-BR-3 cells, the rate of internalization was inversely related to the PEG-PE content of the immunoliposomes.

To study the intracellular disposition of immunoliposomes and their contents, electron microscopy was performed using immunoliposomes containing encapsulated colloidal gold particles. SK-BR-3 cells treated with conventional anti-p185<sup>HER2</sup> immunoliposomes for 30 minutes showed numerous gold-containing



immunoliposomes bound at the cell surface and present intracellularly. Many immunoliposomes were observed contiguous with, and apparently bound to, the cell membrane. Some immunoliposomes were found in coated pits as well as within coated vesicles, endosomes, multivesicular bodies, and lysosomes. This intracellular distribution is consistent with internalization via the coated pit pathway. However, also observed were immunoliposomes appearing to fuse with the cell membrane, without coated pit formation. In addition, some gold particles appeared free within the cytoplasm, not associated with a liposomal capsule or a membrane-bound organelle. Gold particles free within the cytoplasm might have resulted from fusion events between immunoliposomes and the cell membrane. Alternatively, they may have arisen following endocytosis, with escape of the encapsulated gold particles occurring somewhere along the coated pit pathway.

#### Example 6

##### **15 In Vivo Tumor Localization and Biodistribution of Immunoliposomes and Immunoliposome Contents**

Young (4-6 week) female SCID mice were injected with BT-474 cells in the subcutaneous tissue of the flank or of the mammary fat pad, and additionally treated with subcutaneously implanted estrogen pellets to support the tumorigenic growth of these cells. When palpable tumors achieved a size of at least 300 mm<sup>3</sup> (typically 14 days post-inoculation), immunoliposomes were administered by single intravenous injection (via tail vein) or single intraperitoneal injection, in a volume of approximately 200  $\mu$ L containing approximately 1  $\mu$ mole of total lipid. Animals were sacrificed at designated times post-injection, organs perfused with saline *in situ*, and tissues immediately excised for analysis. For biodistribution and imaging analysis after injection of rhodamine-labeled immunoliposomes, freshly excised tissues were fixed and examined by confocal fluorescence microscopy. For quantitative localization of doxorubicin delivered by immunoliposomes, excised tissues were homogenized and subjected to acidified ethanol extraction; the extracted doxorubicin was then measured by

spectrofluorimetric assay as previously described (Gabizon *et al.*, *J. Natl. Cancer Inst.*, 81: 1484-1488 (1989)).

The biodistribution of anti-p185<sup>HER2</sup> immunoliposomes and their ability to localize to and accumulate within tumors *in vivo* was evaluated in a tumor xenograft model. In this model, immunodeficient mice carrying established subcutaneous BT-474 tumor xenografts were treated with immunoliposomes by a single intravenous or intraperitoneal injection. Imaging studies with confocal fluorescence microscopy were performed to detect rhodamine-labeled immunoliposomes within various tissues of animals sacrificed after treatment. Within 6 hours of intravenous injection, rhodamine fluorescence was visualized within the xenografted tumor, while no significant fluorescence was observed within surrounding muscle. This technique, however, did not allow precise delineation of immunoliposomes within tumor tissue. For a quantitative evaluation of the biodistribution and localization of doxorubicin delivered by immunoliposomes, doxorubicin was assayed from tissue extracts of treated animals. 24 hours after intraperitoneal injection, doxorubicin delivered by sterically stabilized anti-p185<sup>HER2</sup> immunoliposomes had accumulated within tumor xenografts, with lower levels of doxorubicin found in surrounding muscle and in blood (Table 2).

Table 2. Anti-P185<sup>HER2</sup> immunoliposome delivery of doxorubicin *in vivo*: biodistribution 24 hrs after single ip injection.

Tissue/Fluid	Doxorubicin Level (% injected dose/g of tissue)
5 Tumor	1.64
Muscle	0.61
Liver	13.46
Ventricular Blood	0.88
10 Tumor/Muscle Ratio	2.69
Tumor/Blood Ratio	1.88

The above examples are provided to illustrate the invention but not to limit its scope. Other variants of the invention will be readily apparent to one of ordinary skill in the art and are encompassed by the appended claims. All  
15 publications, patents, and patent applications cited herein are hereby incorporated by reference.

**WHAT IS CLAIMED IS:**

1. A growth-inhibiting immunoliposome that specifically binds to a cell bearing a characteristic cell surface marker, said immunoliposome comprising:
  - 5 a Fab' domain of an antibody where said Fab' domain specifically binds said marker; and
  - an amphipathic vesicle-forming lipid, said lipid forming a liposome containing no growth-inhibiting therapeutic agent;
  - wherein said immunoliposome inhibits the proliferation of said cell.
- 10 2. The immunoliposome of claim 1, wherein said marker is a growth factor receptor.
3. The immunoliposome of claim 2, wherein said marker is a  
15 growth factor receptor selected from the group consisting of HER1, HER2, HER3 and HER4.
4. The immunoliposome of claim 3, wherein said marker is  
20 HER2.
5. The immunoliposome of claim 4, wherein said Fab' domain is a humanized Fab' domain of an anti-HER2 monoclonal antibody.
6. The immunoliposome of claim 1, wherein said liposome  
25 further comprises a maleimide derivatized phosphatidylethanolamine (M-PE) which forms a thioether linkage to said Fab' domain of an antibody.
7. The immunoliposome of claim 1, wherein said vesicle  
30 forming lipid comprises a lipid selected from the group consisting of a phospholipid, a glycolipid, a sphingolipid, and a sterol.

8. The immunoliposome of claim 1 further comprising a polyethylene glycol derivatized lipid wherein the polyethylene glycol has an average molecular weight of between about 750 D and 5000 D.

5 9. The immunoliposome of claim 1, wherein the average diameter of said liposome ranges from about 50 nm to about 500 nm.

10 10. The immunoliposome of claim 9 wherein the average diameter of said liposome is about 100 nm.

11. The immunoliposome of claim 8 wherein:  
said Fab' domain of an antibody is rhuMAbHER2;  
said Fab' domain of an antibody is conjugated to  
maleimide derivatized phosphatidylethanolamine (M-PE);  
15 said vesicle forming lipid is phosphatidylcholine (PC)  
and cholesterol (Chol); and  
said derivatized lipid is polyethylene glycol derivatized  
phosphatidylethanolamine (PEG-PE) wherein said polyethylene glycol has a  
molecular weight of about 1900 D;  
20 where the ratio PC:Chol:M-PE is about 150:100:3 and said PEG-PE is present in  
an amount up to about 3.6 mole percent of total lipid.

12. A method of inhibiting growth of a cell bearing a  
characteristic cell surface marker, said method comprising contacting said cell with  
25 a growth-inhibiting immunoliposome, said immunoliposome comprising:  
a Fab' domain of an antibody where said Fab' domain  
specifically binds said marker; and  
an amphipathic vesicle-forming lipid, said lipid forming a  
liposome containing no growth-inhibiting therapeutic agent.

13. The method of claim 12, wherein said marker is a growth factor receptor selected from the group consisting of HER1, HER2, HER3 and HER4.

5 14. The method of claim 13, wherein said Fab' domain is a humanized Fab' domain of a monoclonal antibody.

15 15. The method of claim 12, wherein said liposome further comprises a maleimide derivatized phosphatidylethanolamine (M-PE) which forms  
10 a thioether linkage to said antibody.

16. The method of claim 12, wherein said vesicle forming lipid comprises a lipid selected from the group consisting of a phospholipid, a glycolipid, a sphingolipid, and a sterol.

15

17. The method of claim 12, wherein said immunoliposome further comprises a polyethylene glycol derivatized lipid wherein said polyethylene glycol has an average molecular weight of between about 750 D and 5000 D.

20 18. The method of claim 12, wherein the average diameter of said liposome ranges from about 50 nm to about 500 nm.

19. The method of claim 12, wherein:  
said Fab' domain of an antibody is rhuMAbHER2;  
25 said Fab' domain of an antibody is conjugated to maleimide derivatized phosphatidylethanolamine (M-PE);  
said vesicle forming lipid comprises phosphatidylcholine (PC) and cholesterol (Chol); and  
said derivatized lipid is polyethylene glycol derivatized  
30 phosphatidylethanolamine (PEG-PE) wherein said polyethylene glycol has a molecular weight of about 1900 D;

where the ratio PC:Chol:M-DPE is 150:100:3 and said PEG-PE is present in an amount up to 3.6 mole percent of total lipid.

20. An immunoliposome that optimizes internalization of a  
5 therapeutic agent into the cytoplasm of a cell bearing a characteristic cell surface marker, said immunoliposome comprising:

a Fab' domain of an antibody wherein said Fab' domain specifically binds said marker;

an amphipathic vesicle-forming lipid, said vesicle-forming  
10 lipid forming a liposome;

a polyethylene glycol derivatized lipid wherein the polyethylene glycol has an average molecular weight of between about 750 D and 5000 D, the derivatized lipid being present at up to about 3.6 mole percent of the total lipid; and

15 a therapeutic agent contained within said liposome.

21. The immunoliposome of claim 20, wherein said marker is a growth factor receptor selected from the group consisting of HER1, HER2, HER3 and HER4.

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22. The immunoliposome of claim 21, wherein said Fab' domain is a humanized Fab' domain of a monoclonal antibody.

23. The immunoliposome of claim 20, wherein said liposome  
25 further comprises a maleimide derivatized phosphatidylethanolamine (M-PE) which forms said thioether linkage to Fab' domain of an antibody.

24. The immunoliposome of claim 20, wherein said vesicle forming lipid is selected from the group consisting of a phospholipid, a glycolipid,  
30 a sphingolipid, and a sterol.

25. The immunoliposome of claim 20, wherein the average diameter of said liposome ranges from about 50 nm to about 500 nm.

26. The immunoliposome of claim 20, wherein said therapeutic agent is selected from the group consisting of daunomycin, idarubicin, mitoxantrone, mitomycin, cisplatin and other Platinum II analogs, vincristine, epirubicin, aclacinomycin, methotrexate, etoposide, doxorubicin, cytosine arabinoside, fluorouracil and other fluorinated pyrimidines, purines, or nucleosides, bleomycin, mitomycin, plicamycin, dactinomycin, cyclophosphamide and derivatives thereof, thiotepa, BCNU, taxol, taxotere and other taxane derivatives and isolates, camptothecins, polypeptides, a nucleic acid, a nucleic acid having a phosphorothioate internucleotide linkage, and a nucleic acid having a polyamide internucleotide linkage.

27. The immunoliposome of claim 20, wherein:  
said Fab' domain of an antibody is rhuMAbHER2;  
said vesicle forming lipid comprises phosphatidylcholine (PC) and cholesterol (Chol);  
said derivatized lipid is polyethylene glycol derivatized phosphatidylethanolamine (PEG-PE) wherein said polyethylene glycol has a molecular weight of about 1900 D;  
said antibody is conjugated to maleimide derivatized phosphatidylethanolamine (M-PE);  
wherein said ratio of PC:Chol:M-DPE is 150:100:3:6 and  
said PEG-PE is present in an amount less than 3.6 mole percent of total lipid.

28. A method of optimizing internalization of a therapeutic agent into the cytoplasm of a cell bearing a characteristic cell surface marker, said method comprising the steps of  
a) contacting said cell with an immunoliposome, said immunoliposome comprising:



i) an Fab' domain of an antibody where said Fab' domain specifically binds said marker;

ii) an amphipathic vesicle-forming lipid, said lipid forming a liposome;

5                   iii) a polyethylene glycol derivatized lipid wherein the polyethylene glycol has an average molecular weight of between about 750 D and 5000 D, the derivatized lipid being present at less than 3.6 mole percent of the total lipid; and

iv) a therapeutic agent contained within said liposome;

10    and

b) internalizing said liposome itself into said cell.

29.    The method of claim 28, wherein said internalizing comprises transporting said liposome directly into the cytoplasm of said cell.

15

30.    The method of claim 28, wherein said marker is a growth factor receptor selected from the group consisting of HER1, HER2, HER3 and HER4.

20                   31.    The method of claim 30, wherein said Fab' domain is a humanized Fab' domain of a monoclonal antibody.

32.    The method of claim 28, wherein said vesicle forming lipid is selected from the group consisting of a phospholipid, a glycolipid, a sphingolipid, and a sterol.

25                   33.    The immunoliposome of claim 28, wherein said therapeutic agent is selected from the group consisting of daunomycin, idarubicin, mitoxantrone, mitomycin, cisplatin and other Platinum II analogs, vincristine, 30    epirubicin, aclacinomycin, methotrexate, etoposide, doxorubicin, cytosine arabinoside, fluorouracil and other fluorinated pyrimidines, purines, or nucleosides,

bleomycin, mitomycin, plicamycin, dactinomycin, cyclophosphamide and derivatives thereof, thiotepa, BCNU, taxol, taxotere and other taxane derivatives and isolates, camptothecins, polypeptides, a nucleic acid, a nucleic acid having a phosphorothioate internucleotide linkage, and a nucleic acid having a polyamide  
5 internucleotide linkage.

34. The immunoliposome of claim 28, wherein:

said Fab' domain of an antibody is rhuMAbHER2;

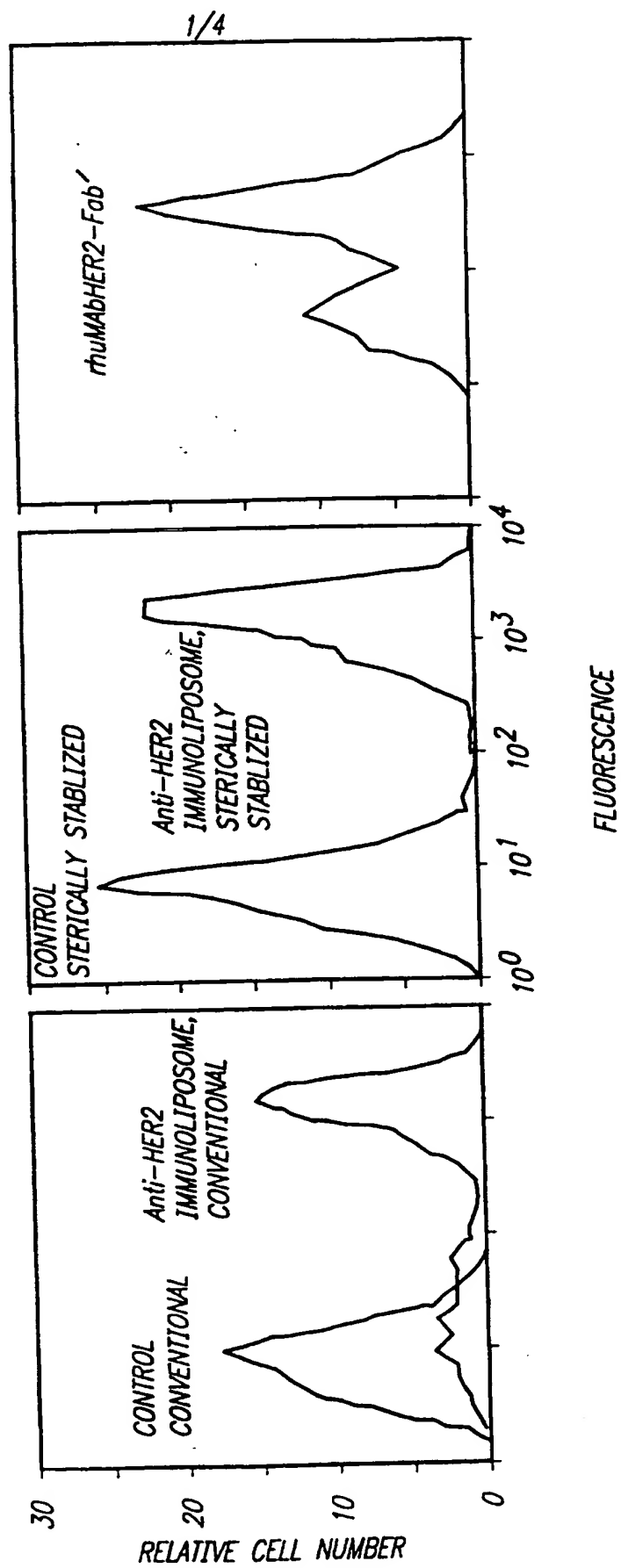
said vesicle forming lipid comprises  
10 phosphatidylcholine (PC) and cholesterol (Chol);

said derivatized lipid is polyethylene glycol derivatized phosphatidylethanolamine (PEG-PE) wherein said polyethylene glycol has a molecular weight of about 1900 D;

said Fab' domain of an antibody is conjugated to  
15 maleimide derivatized phosphatidylethanolamine (M-PE);

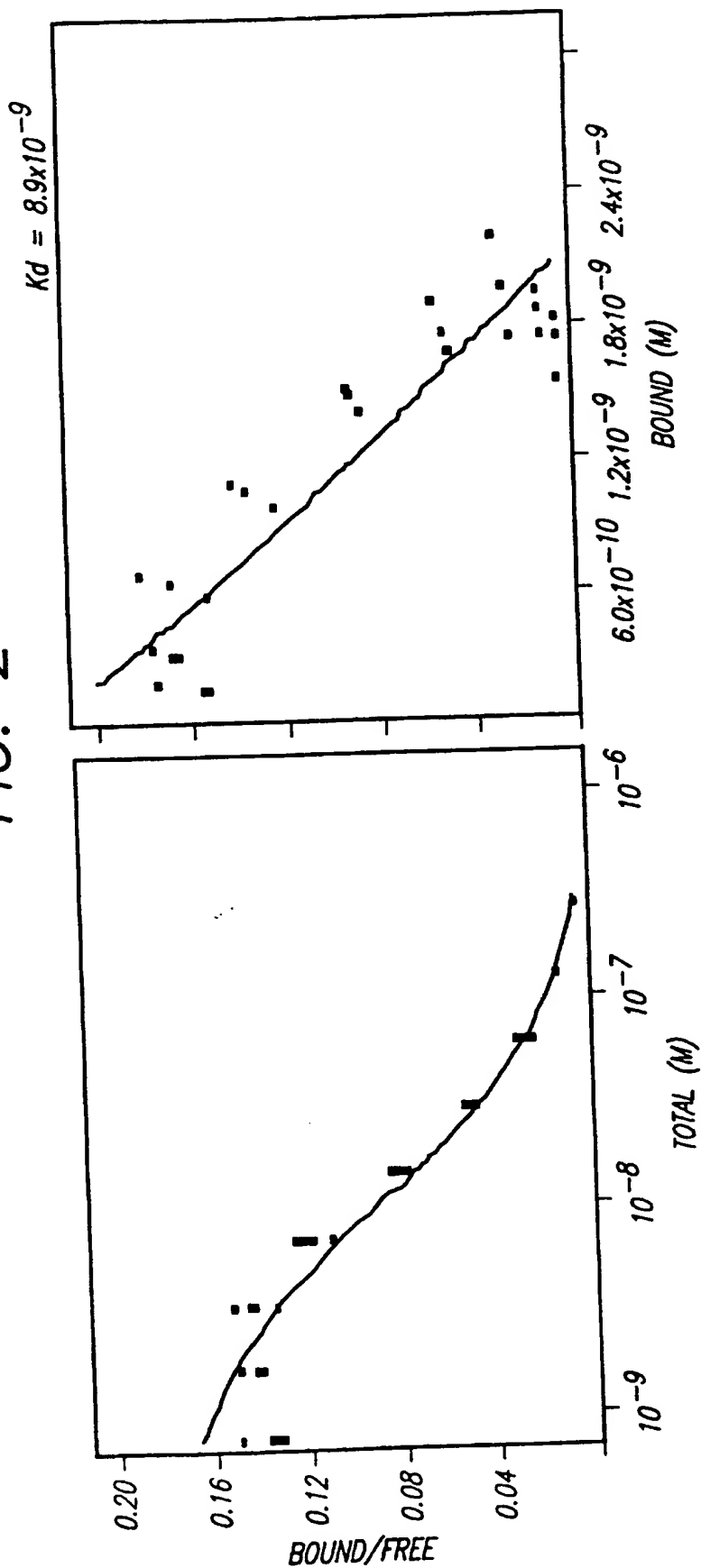
wherein said ratio of PC:Chol:M-DPE is 150:100:3:6  
and said PEG-PE is present in an amount up to 3.6 mole percent of total lipid.

FIG. 1



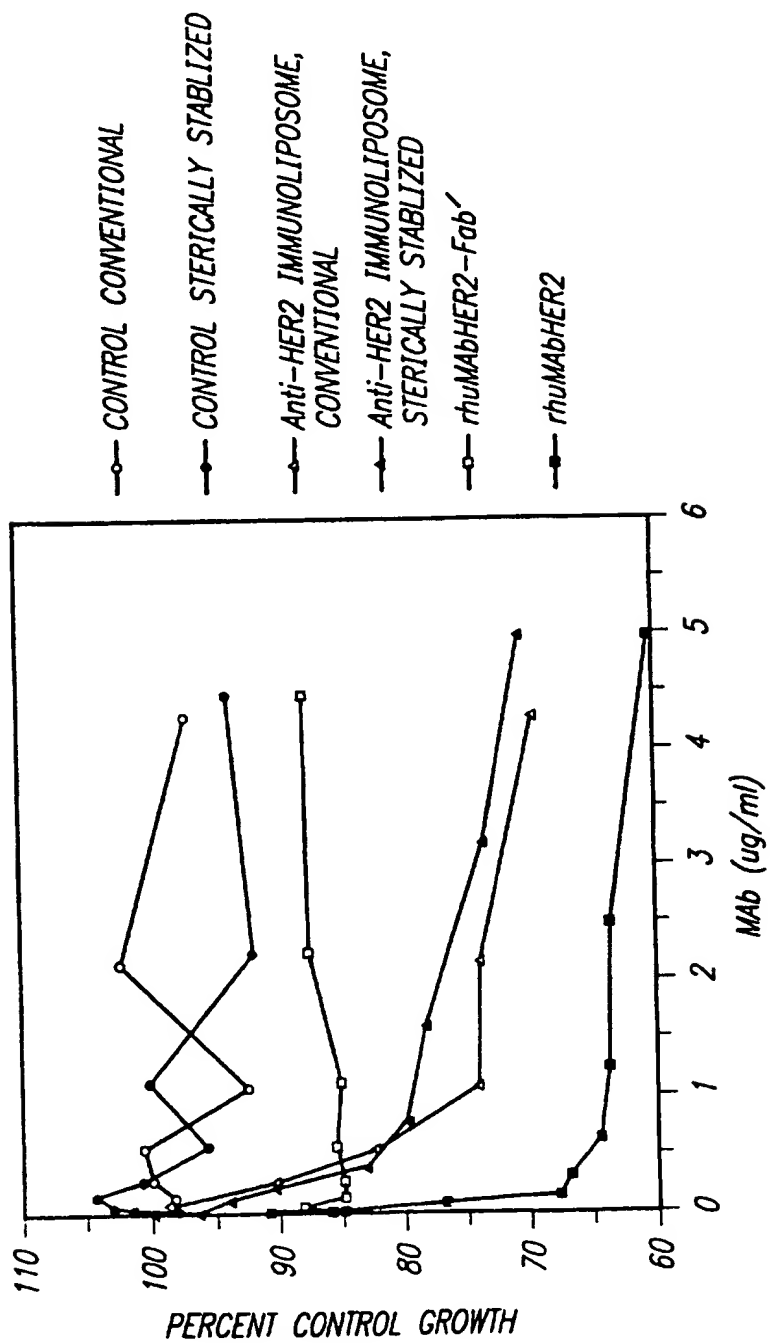
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FIG. 2



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FIG. 3



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FIG. 4A

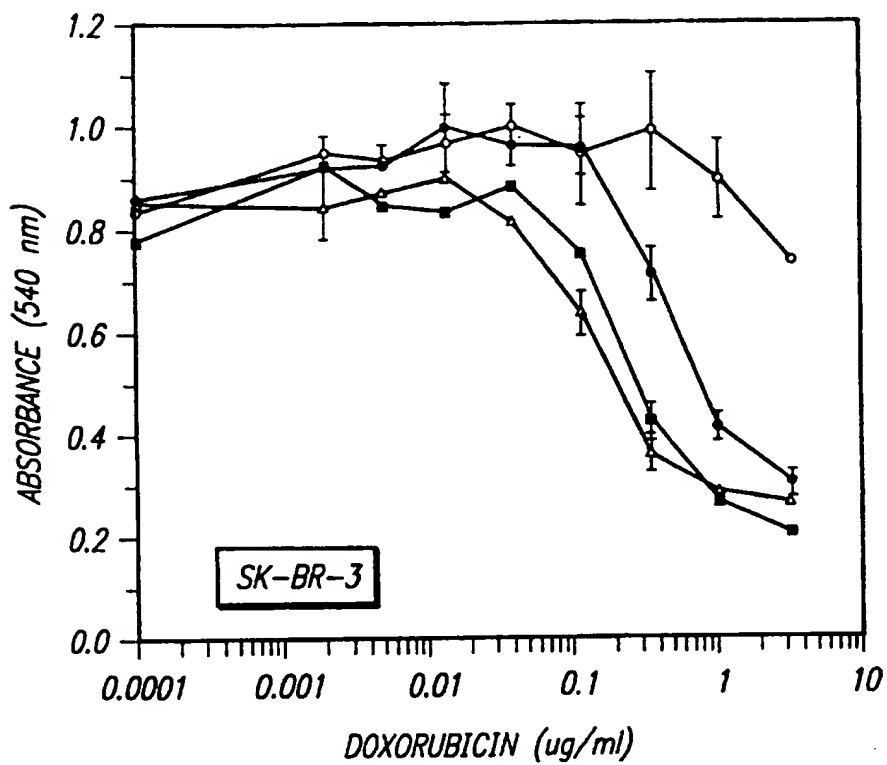
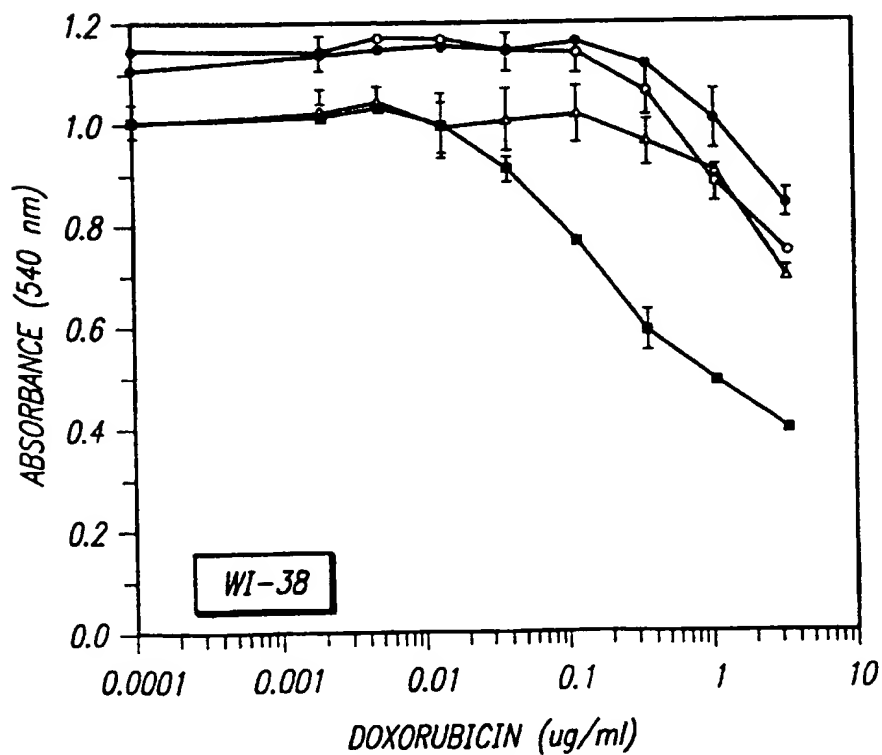


FIG. 4B



## INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US95/14710

## A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) : A61K 39/00, 39/44, 9/127; C07K 16/30, 16/46

US CL : 424/178.1, 450, 812; 530/387.3, 388.8

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 424/178.1, 450, 812; 530/387.3, 388.8

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

APS/ DIALOG (EMBASE, MEDLINE, BIOSYS, LIFESCI, WPI)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	Cancer Research, Volume 52, issued 01 October 1992, S.K. Huang et al., "Microscopic localization of sterically stabilized liposomes in colon carcinoma-bearing mice", pages 5135-5143, see entire document.	1-34
Y	Proceedings of the National Academy of Sciences USA, Volume 89, issued May 1992, P. Carter et al., "Humanization of an anti-p185 <sup>her2</sup> antibody for human cancer therapy", pages 4285-4289, see entire document.	1-34
Y	The Journal of Biological Chemistry, Volume 257, Number 1, issued 10 January 1992, F.J. Martin et al., "Irreversible coupling of immunoglobulin fragments to preformed vesicles", pages 286-288, see entire document.	1-34

☐

Further documents are listed in the continuation of Box C.

☐

See patent family annex.

•	Special categories of cited documents:	T	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
*A	document defining the general state of the art which is not considered to be of particular relevance		
*E	earlier document published on or after the international filing date	*X	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
*L	document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*Y	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
*O	document referring to an oral disclosure, use, exhibition or other means		
*P	document published prior to the international filing date but later than the priority date claimed	*A	document member of the same patent family

Date of the actual completion of the international search

11 JANUARY 1996

Date of mailing of the international search report

06 FEB 1996

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